

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of Brian A. Rosenfeld, M.D. and Michael Breslow

Serial No.: Not yet assigned

Group Art Unit:

Filed: Herewith

Examiner:

For: **SYSTEM AND METHOD FOR PROVIDING CONTINUOUS, EXPERT NETWORK  
CRITICAL CARE SERVICES FROM A REMOTE LOCATION(S)**

Assistant Commissioner for Patents  
Box PATENT APPLICATION  
Washington, D.C. 20231

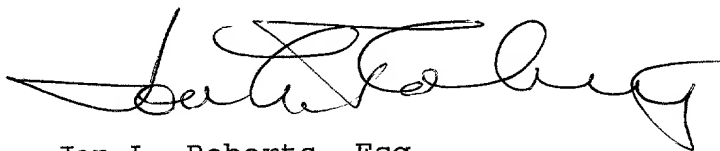
Dear Sir:

Enclosed please find the following:

1. Specification, abstract and claims (2 independent, 12 dependent, 14 total) (111 pages);
2. Informal drawings (57 figures, 56 sheets);
3. Declaration and Power of Attorney;
4. Verified Statement Claiming Small Entity Status - Independent Inventor;
5. Verified Statement Claiming Small Entity Status - Small Business Concern;
6. One check in the amount of \$380.00; and,
7. Certificate of Express mailing.

The Commissioner is hereby authorized to charge any fee deficiency, or credit any overpayment, to Deposit Account No. 18-1579. The Commissioner is also authorized to charge Deposit Account No. 18-1579 for any future fees connected in any way to this application. Two copies of this letter are enclosed.

Respectfully submitted,



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<b>VERIFIED STATEMENT CLAIMING SMALL ENTITY STATUS</b> <b>(37 CFR 1.9(f) &amp; 1.27(c))—SMALL BUSINESS CONCERN</b>	<b>Docket Number (Optional)</b>
Applicant or Patentee: <u>Brian A. Rosenfeld, M.D. and Michael Breslow</u> Serial or Patent No.: <u>Not Yet Issued</u> Filed or Issued: <u>Herewith</u> Title: <u>System and Method For Providing Continuous, Expert Network Critical Care Services From A Remote Location</u>	
I hereby declare that I am <input type="checkbox"/> the owner of the small business concern identified below; <input checked="" type="checkbox"/> an official of the small business concern empowered to act on behalf of the concern identified below:	
NAME OF SMALL BUSINESS CONCERN * <u>ICUSA</u>	
ADDRESS OF SMALL BUSINESS CONCERN <u>2400 Boston Street, Suite 302, Baltimore, MD 21224</u>	
<p>I hereby declare that the above identified small business concern qualifies as a small business concern as defined in 13 CFR 121.12, and reproduced in 37 CFR 1.9(d), for purposes of paying reduced fees to the United States Patent and Trademark Office, in that the number of employees of the concern, including those of its affiliates, does not exceed 500 persons. For purposes of this statement, (1) the number of employees of the business concern is the average over the previous fiscal year of the concern of the persons employed on a full-time, part-time or temporary basis during each of the pay periods of the fiscal year, and (2) concerns are affiliates of each other when either, directly or indirectly, one concern controls or has the power to control the other, or a third party or parties controls or has the power to control both.</p> <p>I hereby declare that rights under contract or law have been conveyed to and remain with the small business concern identified above with regard to the invention described in:</p> <p><input checked="" type="checkbox"/> the specification filed herewith with title as listed above.  <input type="checkbox"/> the application identified above.  <input type="checkbox"/> the patent identified above.</p> <p>If the rights held by the above identified small business concern are not exclusive, each individual, concern or organization having rights in the invention must file separate verified statements averring to their status as small entities, and no rights to the invention are held by any person, other than the inventor, who would not qualify as an independent inventor under 37 CFR 1.9(c) if that person made the invention, or by any concern which would not qualify as a small business concern under 37 CFR 1.9(d), or a nonprofit organization under 37 CFR 1.9(e).</p> <p>Each person, concern, or organization having any rights in the invention is listed below:</p> <p><input type="checkbox"/> no such person, concern or organization exists.  <input checked="" type="checkbox"/> each such person, concern or organization is listed below.</p> <p>Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27).</p> <p>I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlements to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate (37 CFR 1.289(b)).</p> <p>I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.</p>	
NAME OF PERSON SIGNING <u>Robert Pfotenhauer</u>	
TITLE OF PERSON IF OTHER THAN OWNER <u>CEO</u>	
ADDRESS OF PERSON SIGNING <u>2400 Boston Street, Suite 302, Baltimore, MD 21224</u>	
SIGNATURE <u>Robert Pfotenhauer</u>	DATE <u>11/16/99</u>

PTO/SB/09 (11.90)

<b>VERIFIED STATEMENT CLAIMING SMALL ENTITY STATUS (37 CFR 1.9(f) &amp; 1.27(b))—INDEPENDENT INVENTOR</b>	<b>Docket Number (Optional)</b>
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Applicant or Patentee: Brian A. Rosenfeld, M.D. and Michael BreslowSerial or Patent No.: Not Yet IssuedFiled or Issued: HerewithTitle: System and Method for Providing Continuous, Expert Network Critical Care  
Services From A Remote Location(s)

As a below named inventor, I hereby declare that I qualify as an independent inventor as defined in 37 CFR 1.9(c) for purposes of paying reduced fees to the Patent and Trademark Office described in:

- ☐ the specification filed herewith with title as listed above.
- ☒ the application identified above.
- ☐ the patent identified above.

I have not assigned, granted, conveyed or licensed and am under no obligation under convey or law to assign, grant, convey or license, any rights in the invention to any person who would not qualify as an independent inventor under 37 CFR 1.9(c) if that had made the invention, or to any concern which would not qualify as a business concern under 37 CFR 1.9(d) or a nonprofit organization under 37 CFR 1.9(c).

Each person, concern or organization to which I have assigned, granted, conveyed, or licensed or am under an obligation under contract or law to assign, grant, convey, or license any rights in the invention is listed below:

- ☐ No such person, concern, or organization exists.
- ☒ Each such person, concern or organization is listed below.

**ICSUA**

Separate verified statements are required from each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

I acknowledge the duty to file in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b))

I hereby declare that all statements made herein of my own knowledge are true and that all statements made an information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Brian A. Rosenfeld, M.D.

NAME OF INVENTOR

Signature of inventor

Date

Michael Breslow

NAME OF INVENTOR

Signature of inventor

Date

NAME OF INVENTOR

Signature of inventor

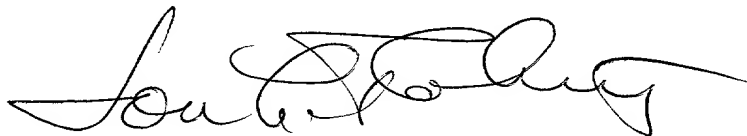
Date

CERTIFICATE OF EXPRESS MAILING

Express Mail Mailing Label Number EL449836293US

Date of Deposit: November 18, 1999

I hereby certify that the patent application of Brian A. Rosenfeld, M.D. and Michael Breslow for a **SYSTEM AND METHOD FOR PROVIDING CONTINUOUS, EXPERT NETWORK CRITICAL CARE SERVICES FROM A REMOTE LOCATION(S)** including the specification, abstract, and claims (2 independent, 12 dependent, 14 total) (1~~1~~<sup>8</sup> pages); informal drawings (57 figures, 56 sheets); declaration and power of attorney; Verified Statement Claiming Small Entity Status - Small Business Concern; Verified Statement Claiming Small Entity status - Independent Inventor; and a check in the amount of \$380.00, are being deposited with the United States Postal Service for "Express Mail" service under 37 C.F.R. § 1.10 on the date indicated above and are addressed to the Assistant Commissioner for Patents, Box Patent Application, Washington, D.C. 20231.



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**Title: System and Method for Providing Continuous, Expert Network Critical Care Services from a Remote Location(s)**

**Inventor: Brian A. Rosenfeld, M.D. and Michael Breslow**

**Field of the Invention**

This invention relates generally to the care of patients in Intensive Care Units (ICUs). More particularly this invention is a system and method for care of the critically ill that combines a real-time, multi-node telemedicine network and an integrated, computerized patient care management system to enable specially-trained Intensivists to provide 24-hour/7-day-per-week patient monitoring and management to multiple, geographically dispersed ICUs from both on-site and remote locations.

**Background of the Invention:**

While the severity of illness of ICU patients over the past 15 years has increased dramatically, the level of and type of physician coverage in most ICUs has remained constant. Most ICU patients receive brief minutes of attention during morning rounds from physicians with limited critical care experience. During the remainder of the day and night, nurses are the primary caregivers, with specialists called only **after** patient conditions have started to deteriorate. The result of this mismatch between severity of illness and physician coverage is an unacceptably high ICU mortality rate (10% nationwide), and a high prevalence of avoidable errors that result in clinical complications. In 1998, an Institute of Medicine Roundtable determined that avoidable patient complications were the single largest problem in medical care delivery. In another prominent 1998 study of 1000 patients, 46% experienced an avoidable adverse event in care, with 40% of these errors resulting in serious disability or death.

1 The physicians who can remedy this situation are in critically short supply. Numerous  
2 studies have shown that Intensivists (physicians who have trained and board certified in Critical  
3 Care Medicine) can markedly improve patient outcomes. However, only one-third of all ICU  
4 patients ever has an Intensivist involved in their care, and the number of Intensivists would  
5 need to increase tenfold (nationally) to provide 24-hour coverage to all ICU patients. With the  
6 rapid aging of the population, this shortfall of expertise is going to increase dramatically.

7 Even where Intensivists are present (and especially where they are not), patients suffer  
8 from unnecessary variation in practice. There is little incentive for physicians to develop and  
9 conform to evidence-based best practices (it takes significant work and a change in behavior to  
10 develop and implement them). This variation contributes to sub-optimal outcomes, in both the  
11 quality and cost of care delivered to ICU patients.

12 What is needed is a redesigning of the critical care regimen offered to patients in an  
13 ICU. Rather than the consultative model where a periodic visit takes place and the doctor then  
14 goes away, a more active 24-hour intensivist managed care is required. Further, technology  
15 that leverages the intensivists' expertise and standardizes the care afforded to patients in an ICU  
16 is required. Further, continuous feedback to improve the practice of intensivists in an ICU is  
17 necessary to provide the intervention required to minimize adverse events. This invention  
18 seeks to provide new methods for managing and delivering care to the critically ill.

19 Attempts to automate various aspects of patient care have been the subject of various  
20 inventions. For example, U.S. Patent No. 5,868,669 to Iliff was issued for "Computerized  
21 Medical Diagnostic and Treatment Advice System." The disclosed invention is for a system

1 and method for providing computerized knowledge based medical diagnostic and treatment  
2 advice to the general public over a telephone network.

3 U.S. Patent No. 5,823,948 to Ross, Jr. et al was issued for "Medical Records  
4 Documentation, Tracking and Order Entry System". The disclosed invention is for a system  
5 and method that computerizes medical records, documentation, tracking and order entries. A  
6 teleconferencing system is employed to allow patient and medical personnel to communicate  
7 with each other. A video system can be employed to videotape a patient's consent.

8 U.S. Patent No. 4,878,175 to Norden-Paul et al. was issued for "Method for Generating  
9 Patient-Specific Flowsheets By Adding/Deleting Parameters." The disclosed invention is for  
10 an automated clinical records system for automated entry of bedside equipment results, such as  
11 an EKG monitor, respirator, etc. The system allows for information to be entered at the  
12 bedside using a terminal having input means and a video display.

13 U.S. Patent No. 5,544,649 to David et al. was issued for "Ambulatory Patient Health  
14 Monitoring Techniques Utilizing Interactive Visual Communications." The disclosed invention  
15 is for an interactive visual system, which allows monitoring of patients at remote sites, such as  
16 the patient's home. Electronic equipment and sensors are used at the remote site to obtain data  
17 from the patient, which is sent to the monitoring site. The monitoring site can display and save  
18 the video, audio and patient's data.

19 U.S. Patent No. 5,867,821 to Ballantyne et al. was issued for "Method and Apparatus  
20 for Electronically Accessing and Distributing Personal Health Care Information and Services in  
21 Hospitals and Homes." The disclosed invention is for an automated system and method for

1 distribution and administration of medical services, entertainment services, and electronic  
2 health records for health care facilities.

3 U.S. Patent No. 5,832,450 to Myers et al. issued for "Electronic Medical Record Using  
4 Text Database." The disclosed invention is for an electronic medical record system, which  
5 stores data about patient encounters arising from a content generator in freeform text.

6 U.S. Patent No. 5,812,983 to Kumagai was issued for "Computer Medical File and  
7 Chart System." The disclosed invention is for a system and method which integrates and  
8 displays medical data in which a computer program links a flow sheet of a medical record to  
9 medical charts.

10 U.S. Patent No. 4,489,387 to Lamb et al. was issued for "Method and Apparatus for  
11 Coordinating Medical Procedures." The disclosed invention is for a method and apparatus that  
12 coordinates two or more medical teams to evaluate and treat a patient at the same time without  
13 repeating the same steps.

14 U.S. Patent No. 4,731,725 to Suto et al. issued for "Data Processing System which  
15 Suggests a Pattern of Medical Tests to Reduce the Number of Tests Necessary to Confirm or  
16 Deny a Diagnosis." The disclosed invention is for a data processing system that uses decision  
17 trees for diagnosing a patient's symptoms to confirm or deny the patient's ailment.

18 U.S. Patent No. 5,255,187 to Sorensen issued for "Computer Aided Medical Diagnostic  
19 Method and Apparatus." The disclosed invention is for an interactive computerized diagnostic

1 system which relies on color codes which signify the presence or absence of the possibility of a  
2 disease based on the symptoms a physician provides the system.

3 U.S. Patent No. 5,839,438 to Chen et al. issued for "Intelligent Remote Visual  
4 Monitoring System for Home Health Care Service." The disclosed invention is for a computer-  
5 based remote visual monitoring system, which provides in-home patient health care from a  
6 remote location via ordinary telephone lines.

7 U.S. Patent No. 5,842,978 to Levy was issued for "Supplemental Audio Visual  
8 Emergency Reviewing Apparatus and Method." The disclosed invention is for a system which  
9 videotapes a patient and superimposes the patient's vital statistics onto the videotape.

10 While these invention provide useful records management and diagnostic tool, none of  
11 them provides a comprehensive method for monitoring and providing real time critical care at  
12 disparate ICU's. In short, they are NOT designed for critical care. Further, none of these  
13 inventions provide for the care of a full time intensivist backed by appropriate database and  
14 decision support assistance in the intensive care environment. What would be useful is a system  
15 and method for providing care for the critically ill that maximizes the presence of an intensivist  
16 trained in the care of the critically. Further such a system would standardize the care in ICU's at  
17 a high level and reduce the mortality rate of patients being cared for in ICU's

18 **Summary of the Invention:**

19 The present invention provides a core business of Continuous Expert Care Network  
20 (CXCN) solution for hospital intensive care units (ICUs). This e-solution uses network,

1 database, and decision support technologies to provide 24-hour connectivity between Intensivists  
2 and ICUs. The improved access to clinical information and continuous expert oversight leads to  
3 reduced clinical complications, fewer medical errors, reduced mortality, reduced length of stay,  
4 and reduced overall cost per case.

5 The technology of the present invention as explained below can be implemented all at  
6 once or in stages. Thus the technology, as more fully explained below is available in separate  
7 components to allow for the fact that hospitals may not be able to implement all of the  
8 technology at once. Thus modular pieces (e.g. videoconferencing, vital sign monitoring with  
9 smart alarms, hand-held physician productivity tools, etc.) can be implemented, all of which can  
10 add value in a stand-alone capacity. First amongst these offerings will be an Intensivist Decision  
11 Support System, a stand-alone software application that codifies evidence-based, best practice  
12 medicine for 150 common ICU clinical scenarios. These support algorithms are explained more  
13 fully below.

14 The "Command Center" model, again as more fully set forth below, will ultimately give  
15 way to a more distributed remote management model where Intensivists and other physicians can  
16 access ICU patients and clinicians (voice, video, data) from their office or home. In this  
17 scenario, the present invention will be available in hospital applications that centralize ICU  
18 information, and offer physicians web-based applications that provide them with real-time  
19 connectivity to this information and to the ICUs. This access and connectivity will enable  
20 physicians to monitor and care for their patients remotely. These products will be natural  
21 extensions and adaptations of the present invention and the existing applications disclosed herein

that those skilled in the art will appreciate and which do not depart from the scope of the invention as disclosed herein.

The present invention addresses these issues and shortcomings of the existing situation in intensive care, and its shortfalls via two major thrusts. First, an integrated video/voice/data network application enables continuous real-time management of ICU patients from a remote setting. Second, a client-server database application B integrated to the remote care network B provides the data analysis, data presentation, productivity tools and expert knowledge base that enables a single Intensivist to manage the care of up to 40 patients simultaneously. The combination of these two thrusts B care management from a remote location and new, technology-enhanced efficiency of Intensivist efforts B allows health care systems to economically raise the standard of care in their ICUs to one of 24x7 continuous Intensivist oversight.

It is therefore an object of the present invention to reduce avoidable complications in an ICU.

It is a further object of the present invention to reduce unexplained variations in resource utilization in an ICU.

It is a further objective of the present invention to mitigate the serious shortage of intensivists.

It is yet another objective of the present invention to reduce the occurrence of adverse events in an ICU.

1 It is a further objective of the present invention to standardize the care at a high level  
2 among ICUs.

3 It is yet another objective of the present invention to reduce the cost of ICU care.

4 It is yet another objective of the present invention to dramatically decrease the mortality  
5 in an ICU.

6 It is yet another objective of the present invention to bring information from the ICU to  
7 the intensivist, rather than bring the intensivist to the ICU.

8 It is a further objective of the present invention to combine tele-medical systems  
9 comprising two-way audio/video communication with a continuous real time feed of clinical  
10 information to enable the intensivist to oversee care within the ICU.

11 It is a further objective of the present invention to allow intensivists to monitor ICUs  
12 from a site remote from each individual ICU.

13 It is a further objective of the present invention to bring organized detailed clinical  
14 information to the intensivist, thereby providing standardized care in the ICU.

15 It is yet another objective of the present invention to utilize knowledge-based software to  
16 use rules, logic, and expertise to provide preliminary analysis and warnings for the intensivists.

17 The present invention comprises a command center/remote location, which is  
18 electronically linked to ICUs remote from the command center/remote location. The command  
19 center/remote location is manned by intensivists 24 hours a day, seven days per week. Each ICU  
20 comprises a nurse's station, to which data flows from individual beds in the ICU. Each patient in  
21 the ICU is monitored by a video camera, as well as by clinical monitors typical for the intensive



care unit. These monitors provide constant real time patient information to the nurse's station, which in turn provides that information over a dedicated T-1 (high bandwidth) line to the ICU command center/remote location. As noted earlier, the command center/remote location is remote from the ICU, thereby allowing the command center/remote location to simultaneously monitor a number of patients in different ICUs remote from the command center/remote location.

At each command center/remote location, video monitors exist so that the intensivist can visually monitor patients within the ICU. Further, the intensivist can steer and zoom the video camera near each patient so that specific views of the patient may be obtained, both up close and generally. Audio links allow intensivists to talk to patients and staff at an ICU bed location and allow those individuals to converse with the intensivist.

Clinical data is constantly monitored and presented to the command center/remote location in real time so that the intensivist can not only monitor the video of the patient but also see the vital signs as transmitted from the bedside. The signals from the clinical data and video data are submitted to a relational database, which comprises 1) standardized guidelines for the care of the critically ill, 2) various algorithms to support the intensive care regimen, 3) order writing software so that knowledge-based recommendations and prescriptions for medication can be made based upon the clinical data, and 4) knowledge-based vital sign/hemodynamic algorithms that key the intensivist to engage in early intervention to minimize adverse events.

The advantage of the present invention is that intensivists see all patients at a plurality of ICU's at all times. Further, there is a continuous proactive intensivist care of all patients within

1 the ICU, thereby minimizing adverse events. Intervention is triggered by evidence-based data-  
2 driven feedback to the intensivist so that standardized care can be provided across a plurality of  
3 ICUs.

4 The economic benefits of the present invention are manifold. For the first time, 24-hour  
5 a day, seven day a week intensivist care for patients in an ICU can be obtained. Further, more  
6 timely interventions in the care of the patients can be created by the knowledge-based guidelines  
7 of the present invention, thereby minimizing complications and adverse events. This in turn will  
8 lead to a reduced mortality within the ICU, and hence, a reduced liability cost due to the  
9 dramatic reduction in avoidable errors in health care.

10 By providing timely interventions, the length of stay within the ICU can be greatly  
11 reduced, thereby allowing more critically ill patients to be cared for in the ICU.

12 In addition, by reviewing and standardizing the care afforded to patients in an ICU, a  
13 more standardized practice across a variety of ICUs can be achieved. This will lead to more  
14 cost-effective care within the ICU, and reduced ancillary cost for the care of the critically ill.

15 The overall architecture of the present invention comprises a "pod." The pod comprises a  
16 tele-medicine command center/remote location connected to a plurality multiple ICUs at various  
17 locations. The connection between the command center/remote location and the ICUs is via a  
18 dedicated wide-area network linking the ICUs to the command center/remote location and a team  
19 of intensivists who integrate their services to provide 24-hour, seven day a week care to all of the  
20 pod ICUs.

1           The pod is connected via a wide-area network using dedicated T-1 lines, for example,  
2   with redundant backup. This network provides reliable, high speed secure transmission of  
3   clinical data and video/audio signals between each patient room and the command center/remote  
4   location. The use of a T-1 line is not meant as a limitation. It is expected that more and higher  
5   bandwidth networks will become available. Such high bandwidth networks would come within  
6   the scope of the invention as well.

7           Each patient room is equipped with a pan/tilt/zoom video camera with audio and speaker  
8   to enable full videoconferencing capability. In addition, computer workstations are dedicated for  
9   exclusive physician use in each ICU, preferably at the nurse's station. Intensivists use the  
10   workstations to view patient information, consult decision support information, record their  
11   notes, and generate patient orders.

12           The patient management software used by intensivists is provided across the pod.  
13   Updates and changes made to the record are available at both the ICU and the command  
14   center/remote location for any given patient.

15           Each command center/remote location contains at least three workstations: one for the  
16   intensivist, one for the critical care registered nurse, and one for a clerk/administrative person.

17           The intensivist workstation comprises separate monitors for displaying ICU video images  
18   of patients and/or ICU personnel, output from bedside monitoring equipment, patient clinical  
19   data comprising history, notes, lab reports, etc., and decision support information. The staff at  
20   the command center/remote location are able to activate and control the cameras in each patient's  
21   room so that appropriate visual views of the patient can be generated.

Intensivists are able to switch between rooms and patients and can monitor at least two rooms simultaneously via the video screens. Patient data such as X-ray and ECG images are scanned and transmitted to the command center/remote location upon request of the intensivist.

Remote patient management is utilized in the present invention's critical care program to supplement traditional onsite care. The rationale underlying the remote patient management of the present invention is that critically ill patients are inherently unstable and require continuous expert care that is not now offered in existing ICU monitoring regimens. Further, remote monitoring allows a single intensivist to care for patients in multiple ICU locations, thereby creating an efficiency that makes continuous care feasible.

Remote intensivist care of the present invention is proactive. Intensivists will order needed therapies and check results of tests and monitor modalities in a more timely fashion than is currently offered. Patients can be observed visually when needed using the ceiling-mounted cameras in each room.

Command center/remote location personnel communicate with ICU staff through videoconferencing and through "hot phones," which are dedicated telephones directly linked between the command center/remote location and the ICU. These communications links are used to discuss patient care issues and to communicate when a new order has been generated.

Intensivists document important events occurring during their shift in progress notes generated on the command center/remote location computer terminal.

Intensivists detect impending problems by intermittently screening patient data, including both real time and continuously stored vital sign data. Patient severity of illness determines the frequency with which each patient's data is reviewed by the intensivists.

#### **Brief Description of the Figures**

Figure 1 illustrates the logical data structure for billing, insurance and demographic information

Figure 1A illustrates the logical data structure for billing, insurance and demographic information (cont)

Figure 2 illustrates the command center logical data structure

Figure 2A illustrates the command center logical data structure (cont)

Figure 3 illustrates the logical data structure for creating a medical history

Figure 4 illustrates the logical data structure for creating notes relating to patient treatment and diagnosis

Figure 4A illustrates the logical data structure for creating notes relating to patient treatment and diagnosis (cont)

Figure 4B illustrates the logical data structure for creating notes relating to patient treatment and diagnosis (cont)

Figure 5 illustrates the logical data structure for entry of medical orders

Figure 6 illustrates the logical data structure for patient care, laboratory testing and diagnostic imaging

1 Figure 6A illustrates the logical data structure for patient care, laboratory testing and  
2 diagnostic imaging (cont)

3 Figure 7 illustrates the logical data structure for categories of information that are  
4 permitted to be presented to intensivists and other care givers by the system

5 Figure 8 illustrates the logical data structure for documenting patient vital signs

6 Figure 8A illustrates the logical data structure for documenting patient vital signs (cont)

7 Figure 9 illustrates the distributed architecture of the present invention

8 Figure 10 illustrates the system architecture of the present invention

9 Figure 11 illustrates the decision support algorithm for decision support algorithm for  
10 diagnosis and treatment of pancreatitis.

11 Figure 12 illustrates the vital signs data flow.

12 Figure 13A illustrates capture and display of diagnostic imaging.

13 Figure 13B illustrates establishing videoconferencing in the present invention.

14 Figure 14 illustrates the physician resources order writing data interface of the present  
15 invention.

16 Figure 15 illustrates the physician resources database data interface of the present  
17 invention.

18 Figure 16 illustrates the automated coding and billing system integrated with the  
19 workflow and dataflow of the present invention.

20 Figure 17 illustrates the order writing data flow of the present invention.

21 Figure 18 illustrates the event log flow of the present invention.



Figure 34A illustrates the wide complex QRS Tachycardia decision support algorithm  
(cont)

Figure 41 illustrates the assessment of sedation decision support algorithm

Figure 41A illustrates the assessment of sedation decision support algorithm (cont)

Figure 42 illustrates the bolus sliding scale midazolam decision support algorithm

Figure 43 illustrates the sedation assessment algorithm decision support algorithm

Figure 44 illustrates the short term sedation process decision support algorithm

Figure 45 illustrates the respiratory isolation decision support algorithm

Figure 47 illustrates the empiric meningitis treatment decision support algorithm

Figure 48 illustrates the ventilator weaning decision support algorithm

Figure 48A illustrates the ventilator weaning decision support algorithm (cont)

Figure 49 illustrates the warfarin dosing decision support algorithm

Figure 51 illustrates the HIT-2 diagnostic decision support algorithm

## **Definitions of Terms and Data**

In the following Detailed description of the Invention, a number of modules and procedures are described. For purposes of definitions, the following module definitions apply and are more fully amplified in the descriptions of the figures that follow:

### **Term Definitions:**

Following are a series of definitions for certain terms used in this specification:

Insurance carrier: This is a table of all the valid insurance carriers listed in the system of the present invention.



1 Patient guarantor: Provides the insurance guarantor information for a given patient.

2 Patient information: Provides demographic information for each patient.

3 Medical event date history: This contains the various disorders of the patient and the  
4 dates associated with major medical events relating to those disorders.

5 Medical history: Contains non-major system medical history of a patient.

6 Drug: Contains what medication and allergies have been identified for a patient at  
7 admission.

8 Address: Contains the address or addresses for a given patient.

9 Patient visit: There may be multiple records for any given patient, since the patient may  
10 visit the ICU on more than one occasion. This file contains a record of each visit to an ICU by a  
11 patient.

12 Physician-patient task: Contains the task that had been defined for each patient.

13 Present illness: This contains a textual description of the patient illness for the specific  
14 ICU visit.

15 Physical exam: This contains the information gathered as a result of a physical  
16 examination of the patient during the admission to the ICU.

17 Surgical fluids: This provides all the information related to the fluids provided during  
18 surgery.

19 Surgery: This contains all information pertaining to any surgical procedure performed on  
20 a patient while the patient is at the ICU.

1 Patient admit: This provides general information that needs to be gathered when a patient  
2 is admitted into the ICU.

3 Medical orders: This provides the general information for all types of medical orders  
4 associated with a given patient.

5 Daily treatment: This contains the treatment provided for a given patient on a given day.

6 Daily diagnosis: This contains the daily diagnosis for a given patient, which includes  
7 neurological, cardiological, pulmonary, renal, endocrinological, and any other diagnosis that may  
8 be associated with a patient.

9 Vital sign information is also critical to the administration of care in the ICU. A number  
10 of different modules collect information relating to patient vital signs. For example:

11 Patient admit: This provides the general information that needs to be gathered when a  
12 patient is admitted to the ICU.

13 Patient visit: This contains a record of each visit to an ICU by a patient.

14 Patient: Provides demographic information for each patient.

15 Vital sign header: This contains general information related to the vital sign data for the  
16 particular patient.

17 Vital sign: Contains the vital sign data taken at specific intervals for a given patient.

18 Hospital: This contains identifying information for a particular hospital where the care is  
19 given.

20 ICU bed: Contains the association for identifying which beds are in a given ICU.

Command center/remote location definitions and modules have also been created for the present invention to allow for the orderly storage and retrieval and entering of data. For example:

Physician-physician (such as nurses and LPN and the like): Contains the names of all of the physicians and physician extenders for the command center/remote location as well as for ICUs associated with the command center/remote location.

Communication: Contains all of the various types of communication vehicles used to contact an individual physician or physician extender.

Physician role: Contains the role a physician is playing for a given patient, (i.e., primary care, consultant, etc.)

Patient: Provides demographic information for each patient.

Command center/remote location: Provides identifying information for a particular command center/remote location.

Hospital: Contains identifying information for a particular hospital wherein an ICU is located.

ICU: Contains identifying information for an ICU at a hospital.

ICU bed: Contains the association for identifying which beds are in a given hospital.

ICU patient location: Provides the association between an ICU and a patient and identifies where a patient is located within an ICU in a particular hospital.

1 The order entry functionality of the present invention provides a critical service for  
2 obtaining information on the patient during admission, medical orders, and procedures provided  
3 to the patient during the ICU stay. For example:

4 Radiology: Contains all radiology performed on a particular patient.

5 Radiology results: Contains the results of each radiology test performed on the particular  
6 patient.

7 Drugs: Contains all relevant information for all the drugs that a patient has been  
8 administered.

9 Laboratory: Contains all laboratory tests ordered for a patient.

10 Microbiology result: Contains the results of microbiology organisms taken on a patient.

11 Laboratory result: Contains the results for a laboratory test ordered for a particular  
12 patient.

### 13 **Detailed Description of the Invention**

14 The present invention is a system and method for remote monitoring of ICU's from a  
15 distant command center/remote location. By monitoring a plurality of ICU's remotely,  
16 intensivists can better spread their expertise over more ICU beds that heretofore achievable. The  
17 presence of 24-hour a day/7 day-per-week intensivist care dramatically decreases the mortality  
18 rates associated with ICU care.

19 Referring to Figures 1 and 1A, the Billing and Demographic data structure of the present  
20 invention is illustrated. Patient demographic information 9010 is collected on the particular  
21 patient. This information comprises all the typical kinds of information one would normally

gather on a patient such as first name, last name, telephone number, marital status, and other types of information. Patient insurance information **9012** is collected and associated with the patient demographic information **9010**. Patient insurance information **9012** relates to information on the type of accident and related information such as employment, employer name, place of service, and other information that would relate to the accident that actually occurred (if at all) and which would have to be reported to an insurance agency. This information is associated with the patient demographic information which assigns the unique patient ID to the particular patient.

Insurance plan information **9008** is also created and stored and comprises insurance carrier ID's, the plan name, policy number, and group number. This information on the insurance plan **9008** is also associated with the patient ID and demographic information **9010**.

Physician information **9002** is also created and stored for each physician associated with the system of the present invention. Information such as first and last name, credentials, and other information concerning the physician is saved. In addition, the physician's role is identified **9004** and information concerning the physician and the physician's role is associated with the particular patient via the patient ID stored in the demographic information **9010**.

Patient's are entered into the hospital by a hospital representative **9006** who has a representative ID which also is ultimately associated with the patient ID. In addition, communications data **9000** is stored concerning how a representative can be reached (cell phone, home phone etc.).

Referring now to Figure 1A, the Overall Billing and Insurance data structure is illustrated. An insurance provider number **9014** is also stored in the system. Each physician is given a provider number and provider ID by each insurance company. Thus data must be stored

1 regarding the ID that is given to a particular physician by each insurance provider. This  
2 information is also stored and can be associated ultimately with treatment of the patient.

3 Each patient admitted to the hospital and to the ICU has a patient visit ID associated with  
4 the patient **9017**. This visit ID has patient ID information, ICU information, admission date, and  
5 other information relevant to the specific visit. This information is illustrated in Figure 1A. The  
6 visit ID **9017** is associated with the patient ID **9010** so that each visit can be tracked by patient.

7 Insurance carrier information **9018** is stored by the system and is associated with the  
8 insurance plan information **9008** as appropriate. Thus the particular insurance carrier with its  
9 name, address, and other identifying information **9018** is associated with the type of plan **9008**  
10 carried by the patient. The insurance carrier information **9018** together with the insurance plan  
11 information **9008** is associated with the patient via the patient ID information **9010**.

12 Patient address information **9020** and **9022** are collected for each individual patient and  
13 associated with the patient demographic information **9010**. If there is a patient guarantor, this  
14 information is obtained and stored with information on the guarantor **9026**. Such information as  
15 the guarantor's first and last name, date of birth, and other information is stored and is illustrated  
16 in Figure 1A. Further, the guarantor's address **9024** is also collected and ultimately associated  
17 with the patient demographic information **9010**.

18 Referring to Figures 2 and 2A, the Command Center logical data structure is illustrated.  
19 The various information associated with demographic and insurance information is again used to  
20 manage the care and operations of the command center. Therefore, communications information  
21 **9000** is combined with physician and physician extender (i.e. nurse, LPN and the like)  
22 information **9002** and physician role **9004** to be associated with the demographic information  
23 **9010**. The patient visit information **9017** together with this information is associated with the

1 patient's location which has a unique identifier **9030**. Each location ID has patient ID  
 2 information and visit ID information associated with it.

3 Referring now to Figure 2A, the Command Center logical data structure illustration  
 4 continues. Each ICU bed has an associated location ID which comprises hospital ICU  
 5 information, room number, and bed number **9038**. In addition, and as described earlier,  
 6 instrumentation such as cameras are also associated with the particular patient. Therefore the  
 7 camera setting **9040** will have a location ID relating to the ICU bed as well as have camera value  
 8 settings and associated camera identifier information.

9 Each ICU bed **9038** is associated with an ICU **9032**. Each ICU has information  
 10 associated with it that uniquely identifies the ICU as being associated with the particular  
 11 hospital, and having particular phone numbers, fax numbers, work space addresses, and other  
 12 information, that help to identify the ICU.

13 As noted above, each ICU is associated with a hospital **9034**. Each hospital has a unique  
 14 identifier, as well as its own name, address, and other identifying information. Further, since  
 15 each hospital ICU is to be coordinated through a remote command center, information on the  
 16 remote command center **9036** is associated with the hospital information. Each command center  
 17 has a unique ID and has associated address information stored as well.

18 Thus in the Command Center logical data structure, patient ID information **9010** is linked  
 19 to a patient location **9030** which in turn is associated with an ICU bed **9038** each of which beds  
 20 are uniquely associated an ICU **9032** which is associated with a hospital **9034** which in turn has  
 21 the ICU managed by a command center **9036**.

22 An integral part of the system of the present invention is the recording of medical history.  
 23 Referring to Figure 3, the logical relationship among data elements for medial history is

illustrated. Patient visit information **9017** combined with the physician-physician extender information **9002** is combined with specific note-taking information **9042**. The note information comprises the date and time the notes are taken as well as the note type. The note ID is fed information from the medical history item **9044**, which has its own unique medical ID associated with it. This information comprises medical text, category of information, and other information relevant to the medical history. As noted, this information for medical history **9044** is associated with a note ID **9042**, which in turn is associated with the patient visit and physician information **9017** and **9002**.

Referring to Figure 4, 4A, and 4B, the note-keeping logical data structure of the present invention is illustrated. As noted earlier, the note ID **9042** combines information from visit ID, treating physician, and other information relating to the time the note was entered. Other information is associated with the note ID. Referring first to Figure 4, the patient visit information **9017**, is associated with the note ID **9042**. Various procedural information **9046** is kept by the system of the present invention and is associated with the visit ID **9017**. Physicians are able to create free text patient illness notations **9048** and associate them with the note **9042**. Similarly, free text information regarding functioning of the system **9050** is permitted and also associated with notes regarding the particular patient and procedure **9042**.

Specific notes regarding, for example, surgical procedures are also kept. Surgery notes **9054** are associated with a particular note ID and have such information as anesthesia, surgical diagnosis, elective information, and other related surgical information. Surgical fluids **9052** administered during the course of surgery are associated with the surgery information **9054**. Additionally, any surgical complications **9056** are noted and also associated with the surgery



1 which in turn has an associated note ID.

2 Referring now to Figure 4A, the logical data structure for notes and its description is  
3 continued. An assessment plan **9058** is created and associated with the same note ID for the  
4 particular patient. The plan has a free text field that allows a physician to create the appropriate  
5 assessment plan and associate it with a note ID **9042**.

6 Various daily notes are also kept and associated with the individual note ID **9042**. For  
7 example, the daily mental state **9060** is recorded to document the mental state of the patient. The  
8 daily treatment **9062** administered to the patient is associated with the unique note ID. The daily  
9 diagnosis **9068** is also created and associated with unique note ID **9042**.

10 Any unstable conditions are also noted **9070** and records kept of those conditions.  
11 Similarly mortality performance measures (MPM) information **9072** is kept and associated with  
12 the unique note ID. To the extent that any physical exam **9074** is administered, that physical  
13 exam and any free text created by the physician is associated with the unique ID and records  
14 kept. Allergy information **9076** for the particular patient is also created and stored along with the  
15 allergy type, and allergy name. This information is uniquely associated with the note ID.

16 Referring now to Figure 4B, the Logical Data Structure for the Notes Creation and Storage  
17 description is continued. A specific note item record **9078** is also kept and associated with  
18 unique note ID. This note item comprises the principal diagnosis, the chief complaint, the past  
19 history of the patient, the reason for the note, and various other identifications and flags of  
20 information which help in documenting the patient's condition.

21 Any drugs that are administered to the patient, including dosage, type, and number **9086**  
22 is kept and associated with the unique note ID **9042**.

Procedural note items are also documented **9082**. Procedural notes involve the procedural type, the principal diagnosis, the procedural location, procedural indications, and other information of a procedural nature. Procedural description information **9088** is kept as input to the procedural note item. This information is also associated with a procedural evaluation **9084** which comprises text describing the procedural evaluation that occurred. These three items, the procedural description **9088**, procedural evaluation **9084**, and procedural note items **9082**, are all uniquely associated with the note ID **9042**.

Referring now to Figure 5, the Logical Data Structure of the Medical Order Functionality of the Present Invention is illustrated. Each medical order **9092** has a unique order ID associated with it. This information derives its uniqueness from the visit ID, the representative ID, and various information about the date in which the order was created and other such relevant information. Any non-drug orders **9090** are associated with a unique non-drug order ID. The order is classified, identified, and free text can be created by the physician to describe the order. This information in the non-drug order **9090** is associated with the unique medical order for that particular patient **9092**.

Again physician and physician extender identification information **9002** is also uniquely associated with the medical order to identify the physician involved in creating the particular order in question.

Drug orders **9094** are created each with its own unique drug order ID. Various information is collected as part of the drug order including the type of drug, the dosage, start date, frequency, stop date, to name but a few elements typical of a drug order. The drug order information **9094** is associated with the unique medical order ID **9092** assigned to that particular

1 patient. All of the medical order information is associated with patient visit information **9017**  
2 which allows that information to be uniquely identified with a particular patient for a particular  
3 visit.

4 Referring again to Figure 4B, the system is also capable of annotating and storing various  
5 log items **9080**. For example, an event log item is given a number, a patient profile item has its  
6 own number, as do neurological, cardiographic, pulmonary, renal, and other events can have log  
7 items associated with them and may be used as input to any of the note taking of the present  
8 invention.

9 Referring to Figure 6 and 6A, the logical data structure of the patient care functionality of  
10 the present invention is illustrated. Each patient visit with its unique ID **9017** has a number of  
11 other pieced of information associated with it. For example, physician-patient tasks are tracked  
12 **9098** and have a unique task ID associated with them. The patient code status **9096** is  
13 documented and associated with the physician-patient task **9098** task ID. This information is  
14 uniquely associated with the patient visit via the the patient visit ID **9017**.

15 Laboratory information **9100** has a unique lab ID associated with it. That information is  
16 keyed to the visit ID and records the specimen taken, the date it was taken, and various other  
17 information germane to the laboratory procedure involved. Other lab procedures **9102** are also  
18 documented with another unique ID. "Other" lab ID is associated with the laboratory ID **9100**  
19 which again is uniquely associated with the particular patient.

20 Microbiological studies **9104** are documented together with the date and the date taken  
21 and the type of study involved. Any study of microorganisms **9106** is documented with a unique  
22 microorganism ID. Micro sensitivities **9108** which record the sensitivity to microorganisms and

certain antibiotics is recorded and associated with the microorganism ID **9106**. This information in turn is associated with a microbiological study **9104**, all of which is associated with the unique patient visit ID **9107**.

Respiratory studies **9101** are also recorded with unique identification numbers and a description. This information is again associated with the patient visit ID **9017**.

Referring now to Figure 6A, the logical data structure of the patient care functionality of the Present Invention is further illustrated. Other organism studies **9118** are also conducted to determine any other conditions associated with microorganisms that might exist with the particular patient. This other organism information **9118** is associated with the microorganism studies **9106** which in turn is associated with the microbiology category of information of the present invention **9104**.

Various diagnostic imaging also takes place and is recorded. This image information **9114** has unique image ID associated with each image and comprises associated information such as the image type, the date performed, and other information relevant to the diagnostic imagery. The result of the image taken **9116** is also uniquely identified with the image ID and a unique image result ID. This information is associated with the image information **9114** which again is uniquely associated with the patient visit ID.

Various intake and output for the patient's biological functioning is recorded **9110**. Intake and output total **9112** is recorded and uniquely associated with the intake/output identification note **9110**. Intake/output totals **9112** also comprised the weight the total taken in, the total out, and five-day cumulative totals for biological functioning of the particular patient.

Referring to Figure 7, The Logical Data Structure Concern with Reference Information

for the present invention is illustrated. This data structure allows only certain ranges of data to be input by care givers into the system. This is accomplished by having categories of information **9120** each category capable of having only certain values. Similarly, each type of data **9126** associated with each category is only permitted to have certain values. This combination of Category and Type results in a Combined ID **9122** which can be used in combination with certain values **9128** to create a value and combination **9124** that can be presented to a care giver viewing and entering data. This effectively limits errors in data entry by only allowing certain values to be entered for given types of data. For example, if only milligrams of a medication are supposed to be administered, this data structure prevents a care giver from administering kilograms of material since it is not a permitted range of data entry. The “nextkey” function **9027** is the function that keeps track of the ID’s that are given during the administration of the present invention. This function insures that only unique ID’s are given and that no identical ID’s are given to two different patient’s for example.

Referring to Figure 8, the Logical Data Structure of the Vital Signs Functionality of the Present Invention is illustrated. Vital sign header information **9120** is created and uniquely associated with the visit ID for the particular patient. This header information comprises a date-time stamp combined with hospital information, medical reference numbers, and identification of the patient. Vital sign details **9122** are also created and uniquely date-time stamped and associated with the particular visit ID for the patient. This information comprises all manner of vital sign information relating to blood pressure, respiration, and other factors. Vital sign information is associated with the patient visit **9017** and the demographic information concerning the patient **9016**. Such associations of information can be the basis for later studies.

Referring to Figure 8A, Additional Vital Sign Logical Data Structures are illustrated. For example, a vital sign log header **9120** is created using the unique hospital ID and medical record numbers. Other information such a patient name, and date-time stamp are also stored. Vital sign log details **9124** are created and associated with the vital sign log header **9120**. For example, blood pressure measurements, respiration, and other factors are all detailed for a particular hospital ID. It should be noted that all vital sign data is logged in and kept by the systems of the present invention. Where vital sign information is received but cannot be associated with a particular patient, such communications are noted as errors.

Vital sign error details **9126** are also recorded and associated with a particular hospital. Information and the vital sign error detail also comprises heart rate, blood pressure, and other information. This information is associated with a vital sign error header **9130** which is associated with the hospital identifier and the patient first and last name and other information. Various vital sign error codes **9128** exist with the present invention and are used in association with the vital sign error detail **9126**. This information however relates to communications of vital sign data that are deemed “errors” as noted above.

Care Net patient location **9132** is recorded and associated with a particular hospital ID and location ID for the particular patient. Carenet is a proprietary product designation of Hewlett-Packard and is kept by the system of the present invention since it identifies the equipment from which measurements come. The ICU bed information **9038** is associated with the Care Net patient location **9132**.

Referring to **Figure 9**, the distributed architecture of the present invention is shown. In concept, the distributed architecture comprises a headquarters component **200**, a

command center/remote location **202**, and a hospital ICU **204**, which, while represented as a single hospital in this illustration, in the preferred embodiment comprises several hospital ICUs at different locations. The headquarters unit **200** comprises a database server and data warehouse functionality, together with a patient information front end. The patient information front end **206** provides patient specific information to the command center/remote location. The database server/warehouse function **208** comprises the amassed information of a wide variety of patients, in their various conditions, treatments, outcomes, and other information of a statistical nature that will assist clinicians and intensivists in treating patients in the ICU. The headquarters' function also serves to allow centralized creation of decision support algorithms and a wide variety of other treatment information that can be centrally managed and thereby standardized across a variety of command center/remote locations. Further, the database server/data warehousing functionality **208** serves to store information coming from command center/remote locations replicating that data so that, in the event of a catastrophic loss of information at the command center/remote location, the information can be duplicated at the command center/remote location once all systems are up and running.

At the hospital ICU **204**, each patient room **232, 234** has a series of bedside monitors and both video and audio monitoring of each patient in the patient room. Each ICU further has a nurse's station with a video camera and monitor **230** so that videoconferencing can go on between the nurses and doctors at the nursing station and those intensivists at the command center/remote location. The monitoring equipment at the ICU is served by a monitor server **236**, which receives and coordinates the transmission of all bedside monitoring and nurses station communication with the command center/remote location. Finally, each ICU has a patient

1 information front end **228**, which receives and transmits to the command center/remote location  
2 information concerning the identity and other characteristics of the patient.

3 Command center/remote location **202** comprises its own video capture and monitoring  
4 capability **212** in order to allow the intensivists to view the patients and information from the  
5 bedside monitoring as well as to have videoconferencing with the nursing station and with  
6 patients as the need arises. Information from the monitor server **236** at the hospital ICU is served  
7 to an HL7 (the language for transmitting hospital/patient/diagnostic data) gateway **214** to a  
8 database server **222**. In this fashion, information from the bedside monitors can be stored for  
9 current and historical analysis. Monitor front ends **216** and **218** allow technicians and command  
10 center/remote location personnel to monitor the incoming data from the patient rooms in the  
11 ICU. Information from the patient information front end **228** is provided to an application server  
12 **224**, having its own patient information front end **226** for aggregating and assembling  
13 information in the database **222** that is associated with individual patients in the ICU.

14 It is expected that there will be a great deal of concurrent hospital data that is necessary to  
15 the implementation of the present invention. It is therefore expected that there will be a legacy  
16 database system **210** having a front end **220** from which intensivists and command center/remote  
17 location personnel can retrieve legacy database information.

18 Referring to **Figure 10**, a system architecture of one embodiment of the present invention  
19 is illustrated. Headquarters **200** comprises an application server **238**, an NT file server **240**, and  
20 Sun SPARC Enterprise 250 **242** and Enterprise network management system **244**, a Cisco 3600  
21 router **246**, a Cisco 2924 switch **248**, and a hot phone **250**. The application server **238** is  
22 designed to monitor and update those applications used at the command center/remote location.



The NT file server serves to monitor, store, and replicate information coming from the command center/remote locations. The SPARC Enterprise 250 server **242** is a disc storage server, for storing and serving information, such as practice guidelines, algorithms, patient information, and all matter of other information records that must be stored in order to support the present invention. As explained below, the SPARC Enterprise 250 server and other components are such as routers and switches are commonly used in the ICU, the command center/remote location, and the headquarters. For example:

The Cisco 3600 router is a multi-function device that combines dial access, routing, and local area network (LAN) to LAN services, as well as the multi-service integration of voice, video, and data in the same device. This is necessary, since the various command center/remote locations, headquarters, and intensive care units all must integrate and transmit video, audio, and data among the various entities.

The Cisco 7204 is a router which provides high speed LAN interconnect, virtual private networks, and Internet access, all of which is required for providing the communication in the network of the present invention; and

The Cisco 2924 switch is an autosensing fast ethernet switch, allowing networked multimedia and virtual LAN support. Multi-level security is also offered in the switch to prevent unauthorized users from gaining access and altering switch configuration. These components are also identified in the figures (below).

The particular commercial systems named here are given as but some examples of equipment available today. The function of these equipment is the important factor. Other similar or improved equipment can also be utilized.

1           The network management system **244** allows the entire traffic and condition of the  
2 network to be monitored and to allow maintenance to take place. The router **246** and switch **248**  
3 is used for communication with the various command center/remote locations that are served by  
4 the Headquarters component. The Headquarters component interacts via frame relay with the  
5 command center/remote location **202**.

6           Command center/remote location **202** comprises an applications server **262** for the  
7 purpose of running various applications for the intensivists and command center/remote location  
8 staff. The NT file server **264** at the command center/remote location allows patient files,  
9 historical files, algorithms, practice standards, and guidelines, to be served to the clinicians and  
10 intensivists to assist in monitoring the patients. The Sun SPARC Enterprise 250 **266** is used to  
11 for storage purposes as noted above. The Enterprise network management system **268** monitors  
12 the overall health of the network of command center/remote locations and intensive care units as  
13 well as the functionality of the individual pieces of equipment within the command  
14 center/remote location. A Cisco 2924 switch **256** and Cisco 7204 router **258**, combined with the  
15 Cisco 3600 router **260** allows for point to point communication over a T1 line, with a plurality of  
16 intensive care units located remotely from the command center/remote location. Hot phones **252**  
17 and **254** allow communication with the headquarters and the intensive care unit.

18           Intensive care unit **204** comprises a Cisco 2924 switch **272** for the purpose of interfacing  
19 with the various audio-video feeds **274**, **276** from the various patient rooms and the nursing  
20 station. A local work station **280** is connected to a scanner **282** which allows data to be input,  
21 scanned, and communicated via the point to point T1 communications to the command  
22 center/remote location. Further, the workstation **280** provides for textual advice and patient

1 orders to be delivered to the intensive care unit for execution. The intensive care unit also  
2 comprises a laser printer **284** for the printing of patient orders and other information relevant to  
3 the care of intensive care patients.

4 Referring to **Figure 11**, the videoconferencing/surveillance/imaging components of the  
5 present invention are illustrated. The hospital ICU **204** comprises a series of video cameras **290**,  
6 which are located in patient rooms and at the nurse's station. Control for the cameras is provided  
7 through an RS424 to RS232 converter **288**, with instructions for imaging emanating from the  
8 workstation at the command center/remote location **252** through the ICU workstation **280**  
9 through a multi-port serial controller **286**. Video feed from the video cameras **290** is provided to  
10 an audio-video switcher **292**, which in turn provides its output to the multi-port serial controller  
11 **286** for subsequent viewing at the nurse's station and at the command center/remote location. Of  
12 equal importance is a microphone feed from the patient and from the nurses. That microphone  
13 **296** provides its signal to an audio line amplifier **294**, which in turn provides an audio feed to the  
14 audio-video switcher **292**. In this way, a patient can provide information, as can nurses who are  
15 visiting the patient during the course of patient care. It is also important that information of an  
16 audio nature be fed to the intensive care unit, both to the patient rooms and to the nurse's station.

17 To do this, the multi-port serial controller **286** provides an audio signal to a reverse audio  
18 switcher **298**, which in turn provides information to speakers **300** that are located at the nurse's  
19 station as well as at the bedside of the patients. Information to the reverse audio switcher is  
20 provided an audio amplifier **302** from information from a video codec **304**, which in turn is  
21 connected to the workstation at the ICU. As noted earlier, a scanner **282** is provided, so that

1 information can be scanned and provided to the command center/remote location **202** and a hot  
2 telephone **278** communicates with a telephone **252** at the command center/remote location.

3 Referring to **Figure 12** the vital signs data flow is illustrated. The monitoring system at  
4 each ICU bedside comprises a monitoring system for monitoring the vital signs for the patient.  
5 The vital sign monitoring system **450** captures vital sign data **452** and transmits that vital sign  
6 data **454** using the HL7 language (the standard processing language for hospital data and  
7 information). The processor at the ICU processes the vital sign data for transmission and storage  
8 purposes and transmits that information to the remote location. Vital sign data is then loaded  
9 into the data base **458**. The data base for each individual patient is then reviewed and process  
10 rules are applied **460** to the vital sign data. These process rules relate to certain alarming  
11 conditions which, if a certain threshold is reached, provides an alarm to the intensivist on duty.  
12 The vital sign alarm **462** is then displaced to the intensivist who can then take appropriate action.  
13 A typical type of rule processing of the vital sign data might be if blood pressure remains at a  
14 certain low level for an extended period of time, or if heart rate remains high for an extended  
15 period of time. In addition a wide range of other rules are provided which will provide an  
16 audible alarm to the intensivist before a critical situation is reached.

17 In addition to the information being provided to the alarming system for the intensivist,  
18 the vital sign data **464** is also transmitted **466** into a database warehouse **468** comprising vital  
19 sign data **470** from not only the individual patient but from all of the patients being cared for in  
20 the ICU. This database warehouse provides the ability to do data mining for trends that can give  
21 rise to additional process rules and vital sign thresholding. In addition to the transmission of  
22 vital sign data **454** to the remote site, the vital sign data is displayed in real time at the ICU **472**.

Referring to **Figure 13(a)** the diagnostic imaging interaction is illustrated. X-rays for example, are created and transmitted to the command center **472**. Additionally, the information could be ACT scan, MRI, or any other method of medical diagnostic imaging. The x-ray image is captured at the command center **474** where it is stored and in addition displayed on the image monitor **476** for the intensivist to review.

6 Referring to **Figure 13(b)** the interactive video session is illustrated. A video  
7 conferencing session is established **478** regarding a particular patient in an ICU bed. Using the  
8 video cameras in each room and/or at the nurses station at the ICU, the patient and/or the nurse  
9 can be viewed **480**. On the other end of the video conferencing session is the intensivist who can  
10 then both visually and orally communicate with the patient and/or nurse **482**.

Referring to **Figure 14** the physician resources and order writing data interface is illustrated. The user interface **484** allows the physicians to access physician resources **486**. These resources provide guideline for the treatment of the critically ill. In this example the intensivist is requested to enter the antibiotic associated with colitis **488**. The system then generates a request for a fecal leukocyte test **490**. This request is translated into an order writing module **496** which results in the actual order for the test **502**. Since the order needs to be transmitted to the appropriate organization for execution, an appropriate order is generated to the microbiology laboratory **500** in this instance. The order results are then achieved **506** and the completion of the order is reported to the order writing assignment manager **496**. In addition, the order writing module **502** also results in a task list **504** of orders for various other individuals in laboratories. In addition, user interface **484** allows the physician to re-enter the physician resources module at any particular location with results of the tests. These tests are then fed into

- 1 the system to continue with the diagnostic algorithm processing of the patient test results **494**.
- 2 The user interface also allows interaction with the resident data base **498**

0944307 11489  
6687 24460

Referring to **Figure 15** the physician resources database data interface is illustrated. User interface **508** allows the intensivist to interact with the physician resources data base **510**. In this example, resident data base **524** which comprises the identification and background of the resident admitting the patient causes an admission diagnosis **526** to be created. In this example a diagnosis of pancreatitis is illustrated. This diagnosis of pancreatitis **522** alerts the physician resources module **510** which causes an entry for the topic pancreatitis **512**. The diagnosis algorithm for pancreatitis **514** is then retrieved and a request for an Apache II score **516** is requested. The system also requests information for operative data **528** describing what if any operations have taken place with respect to this patient, vital sign data **530**, request for laboratory information **532**, past medical history for the patient **534** and patient demographics **536**. All this information is provided to the Apache II score assignment manager **538** which assigns an Apache II score based upon weighted composite up to twenty five different variables. This Apache II score is provided to the Apache II score request module **516**. If the severity based Apache II score is greater than or equal to eight the diagnostic of the system continue **520**. If the Apache II score is less than eight, the patient is triaged to a none ICU bed **518** since the patient will not necessarily require intensive care thereby saving relatively scarce resources of the ICU for those who are truly critically ill.

Referring to **Figure 16** the automated coding/billing work flow and data flow is illustrated. Clearly ICUs must be paid for the care that they give. At the outset of the visit **540** the user interface **542** allows for the input of ICD 9 diagnosis code information concerning complexity of the case, whether the patient is stable, whether the physician involved is the attending physician or consulting physician and all other manner of information required for billing purposes. In

1 addition, resident data **544** is input such as patient demographics, insurance information,  
2 physician, guarantor, the date that the service is provided. All this information is provided to the  
3 data manager **546** which assembles the required data element for subsequent processing. The  
4 data manager sends the demographic, physician, guarantor, insurance and related information to  
5 a bill generator **548** which begins to assemble of the information to subsequently generate a bill.  
6 Clinical information is provided to the CPT code assignment manager which assigns codes based  
7 upon the scores and user input for bill generation purposes. A history of present illness (HPI)  
8 score **560** is generated along with a review of systems (ROS) score **562**. A PFSH score **564** is  
9 generated along with a score relating to the physical exam **566**. An MPM score **568** which is a  
10 score relating to the severity of the illness is also generated. All of these various scores are  
11 provided to the CPT assignment manager **558**. Periodically information is downloaded for  
12 management reports **556**. Once all of the information for the CPT code assignment is generated  
13 that information is provided to the bill generator **548** which assembles all the data elements  
14 needed to generate an HCFA1500 claim form. The input for the bill generator is then verified  
15 **550** where the physician can disagree with code assignments return progress notes and generally  
16 review the bill. This smart processing of the HCFA1500 claim form allows for fewer mistakes  
17 to be made. If there is any error or additional information that is required, the verification  
18 process fails the proposed claim form and information regarding that failure is provided back to  
19 the resident data for completion of any missing items. Once an invoice has been verified as  
20 having the appropriate information to be submitted the HCFA1500 claim form is generated **554**.  
21 Additional information is written to a billing data file **552** for importation to the patient  
22 accounting system of the present invention.



Referring to **Figure 17** the order writing data flow is illustrated. Order entry user interface **600** allows the intensivist to order procedures and medication to assist the patients in the ICU. For example, the intensivist can order an ECG **604**. Thereafter the order is reviewed and a digital signature relating to the intensivist is supplied **606**. Once reviewed and signed off, the order is approved **607** and sent to the data output system **610**. Thereafter the data output system prints the order to the printer in the ICU **616**. For record keeping purposes the order is exported in the HL7 language to the hospital data system **618**. In addition the data output system adds an item to the data base that will subsequently cause an intensivist to check the ECG results. This notification to the task list is provided to the database **614**. In addition, as part of the database an orders file relating to the specific patient is also kept. The fact that and ECG has been ordered is entered in the orders file for that patient.

In a similar fashion using the order entry user interface **600** the intensivist can order medications **602** for a patient. The medication order then is provided to an order checking system **608**. The order checking system retrieves information from the database **614** relating to allergies of the patient and medication list which includes medications which are already being administered to the patient. This allows for the order checking system to check for drug interactions. Further laboratory data is extracted from the database **614** and the order checking system checks to insure that there will be no adverse impact of the recommended dosage upon the renal function of the patient. Once the order checking system **608** is completed, the order is okayed and provided to the order review and signature module **606**. In this module the digital signature of the intensivist is affixed to the order electronically and the order is approved **607**. Thereafter it is provided to the data output system **610** where again the orders are printed for ICU

1 and **616** and for the hospital data system. In this case, any medications that are ordered are then  
2 provided to the medications list file in the database **614** so that the complete list of all  
3 medications that are being administered to the ICU patient is current.

4 Referring to **Figure 18** the event log is illustrated. The database **620** contains all manner  
5 of notes and data relating to the particular patient that is admitted to the ICU. For example,  
6 admission notes **622** are taken upon admission of the patient and stored in the file that is specific  
7 to that patient. Progress notes **624** are created during the patients stay within the ICU to note the  
8 progress the patient is making giving the various treatments. Procedural notes **626** are also  
9 created by the intensivist to note what procedures have taken place and what if any events have  
10 occurred associated with those procedures. Laboratory data such as positive blood cultures are  
11 also stored in the file **628** in the database **620**. Further x-ray data **630** and abnormal CT Scan  
12 results are stored in the database.

13 The result of these individual files are then provided to an event log manager **632**. For  
14 example, admission notes might contain operations performed. Progress notes **624** might relate  
15 to the operations performed. This information is provided to the event log manager **632**.  
16 Admission information is also input to the event log manager as are a listing of the procedures  
17 administered to the patient. To the extent there are positive blood cultures in the laboratory data  
18 **628** those are provided to the event log manager **632** as are abnormal CT scan results. All of this  
19 information is made available through the user interface **634**. Thus the event log presents in a  
20 single location key clinical information from throughout a patients stay in the ICU. The event  
21 log user interface provides caregivers with a snapshot view of all salient events since admission.  
22 All relevant data on procedures and laboratory tests, etc. are presented chronologically.

Referring to **Figure 19** the smart alarms of the present invention are illustrated. The smart alarm system constantly monitors physiologic data (collected once per minute from the bedside monitors) and all other clinical information stored in the database (labs, medications, etc). The periodicity of the collection of data is stated for illustrative purposes only. It is well within the scope of the present invention to collect physiological data at more frequent time intervals. Thus, monitor **636** provides information in HL7 form to the interface engine **638**. The physiological data is then formatted by the interface engine for storage in the database **640** where all patient information is maintained. The rules engine **642** searches for patterns of data indicative of clinical deterioration.

One family of alarms looks for changes in vital signs over time, using pre-configured thresholds. These thresholds are patient-specific and setting/disease-specific. For example, patients with coronary artery disease can develop myocardial ischemia with relatively minor increases in heart rate. Heart rate thresholds for patients with active ischemia (e.g. those with unstable angina in a coronary care unit) are set to detect an absolute heart rate of 75 beats per minute. In contrast, patients with known coronary artery disease in a surgical ICU have alarms set to detect either an absolute heart rate of 95 beats per minute or a 20% increase in heart rate over the baseline. For this alarm, current heart rate, calculated each minute based on the median value over the preceding 5 minutes, is compared each minute to the baseline value (the median value over the preceding 4 hours). Physiologic alarms can be based on multiple variables. For example, one alarm looks for a simultaneous increase in heart rate of 25% and a decrease in blood pressure of 20%, occurring over a time interval of 2 hours. For this alarm, thresholds were initially selected based on the known association between changes in these two variables and

1 adverse clinical events. Actual patient data were then evaluated to determine the magnitude of  
2 change in each variable that yielded the best balance between sensitivity and specificity. This  
3 process was used to set the final thresholds for the rules engine.

4 Alarms also track additional clinical data in the patient database. One alarm tracks  
5 central venous pressure and urine output, because simultaneous decreases in these two variables  
6 can indicate that a patient is developing hypovolemia. Other rules follow laboratory data (e.g.  
7 looking for need to exclude active bleeding and possibly to administer blood).

8 The purpose of the rules engine is to facilitate detection of impending problems and to  
9 automate problem detection thereby allowing for intervention before a condition reaches a crisis  
10 state.

11 Referring to **Figure 20** the procedural note-line log is illustrated. This log allows  
12 clinicians to evaluate the likelihood that a given procedure might result in further complications.  
13 In this example presented in this Figure 20 a catheter removal is illustrated. When a new  
14 catheter is inserted in a patient **648** a procedural note is created on the procedure note creation  
15 user interface **646**. The note is reviewed and a digital signature is attached to the note to  
16 associate the note with a particular intensivist **654**. The procedure is then approved and is  
17 provided to the data output system **656**. The procedural note is then printed on the printer in the  
18 ICU **658** and is exported in HL7 language to the hospital data system **660**. In addition, this also  
19 triggers a billing event and the data output system provides appropriate output to the billing  
20 module **662** to generate an invoice line item. In addition, the note is stored in the emergency  
21 medical record associated with the patient in the database **664**. In addition, the line log is  
22 updated in the database **664** to show what procedure was administrated to a patient at what time.

1 If there is an existing catheter, that is displayed to the intensivist at the procedure note creation  
2 user interface 646. This would show an existing catheter changed over a wire 650. That  
3 information is provided to the line id module 652 which extracts information from the line log in  
4 the database 664. This information results in a note being created and provided to the note  
5 review and signature module 664. Thus the line log contains, for each patient, relevant  
6 information about all in-dwelling catheters, including type and location of the catheter, insertion  
7 date, the most recent date that the catheter was changed over a wire, and the date the catheter was  
8 removed. This information helps clinicians evaluate the likelihood that a given catheter is  
9 infected and guides its subsequent management of that procedure.

#### 10 Evidence-based Guidelines, Algorithms, and Practice Standards

##### 11 Decision Support Algorithms

12 In order to standardize treatment across ICUs at the highest possible level, decision  
13 support algorithms are used in the present invention. These include textual material describing  
14 the topic, scientific treatments and possible complications. This information is available in real  
15 time to assist in all types of clinical decisions from diagnosis to treatment to triage.

16 All connections among components of the present invention are presently with a high  
17 bandwidth T-1 line although this is not meant as a limitation. It is anticipated that other existing  
18 and future high bandwidth communication capabilities, both wired and wireless, as well as  
19 satellite communications will be suitable for the communications anticipated for the present  
20 invention.

21 As noted earlier, a key objective of the present invention is to standardize care and  
22 treatment across ICUs. This is effective in the present invention by providing decision support to

intensivists as well as information concerning the latest care and practice standards for any given condition. As noted in Table 1 below, a wide variety of conditions is noted. Each of the conditions has an associated guideline of practice standard that can be presented to the intensivist who might be faced with that particular condition in a patient. These guidelines of practice standards can be accessed at the command center/remote location or at the ICU to assist in the treatment of the patient. Thus, the general categories of cardiovascular, endocrinology, general, gastrointestinal, hematology, infectious diseases, neurology, pharmacology, pulmonary, renal, surgery, toxicology, trauma all have guidelines and practice standards associated with them.

**Table 1**

**EVIDENCE-BASED GUIDELINES  
ALGORITHMS & PRACTICE STANDARDS**

**DECISION SUPPORT**

**CARDIOVASCULAR**

BRADYARRHYTHMIAS  
CARDIOGENIC SHOCK  
CARDIO-PULMONARY RESUSCITATION GUIDELINES  
CONGESTIVE HEART FAILURE  
EMERGENCY CARDIAC PACING  
FLUID RESUSCITATION  
HYPERTENSIVE CRISIS  
**IMPLANTABLE CARDIO-DEFIBRILLATORS**  
INTRA-AORTIC BALLOON DEVICES  
MAGNESIUM ADMINISTRATION IN PATIENTS  
MANAGEMENT OF HYPOTENSION, INOTROPES  
MYOCARDIAL INFARCTION  
MI WITH LEFT BUNDLE BRANCH BLOCK  
PA CATHETER GUIDELINES & TROUBLE-SHOOTING

PERMANENT PACEMAKERS & INDICATIONS  
PULMONARY EMBOLISM DIAGNOSIS  
PULMONARY EMBOLISM TREATMENT  
SUPRA-VENTRICULAR TACHYARRHYTHMIAS  
UNSTABLE ANGINA  
VENOUS THROMBOEMBOLISM PROPHYLAXIS  
VENOUS THROMBOSIS: DIAGNOSIS & TREATMENT  
VENTRICULAR ARRHYTHMIAS

### **ENDOCRINOLOGY**

ADRENAL INSUFFICIENCY  
DIABETIC KETOACIDOSIS  
HYPERCALCEMIA: DIAGNOSIS & TREATMENT  
HYPERGLYCEMIA: INSULIN TREATMENT  
STEROID REPLACEMENT STRATEGIES  
THYROID DISEASE

### **GENERAL**

DEALING WITH DIFFICULT PATIENTS AND FAMILIES  
END OF LIFE DECISIONS  
ETHICAL GUIDELINES  
PRESSURE ULCERS  
ORGAN PROCUREMENT GUIDELINES

### **GASTROINTESTINAL**

ANTIBIOTIC ASSOCIATED COLITIS  
HEPATIC ENCEPHALOPATHY  
HEPATIC FAILURE  
MANAGEMENT OF PATIENTS WITH ASCITES  
NUTRITIONAL MANAGEMENT  
ACUTE PANCREATITIS  
UPPER GI BLEEDING: STRESS PROPHYLAXIS  
UPPER GI BLEEDING: NON-VARICEAL  
UPPER GI BLEEDING:VARICEAL

## **HEMATOLOGY**

HEPARIN  
HEPARIN-INDUCED THROMBOCYTOPENIA  
THE BLEEDING PATIENT  
THROMBOCYTOPENIA  
THROMBOLYTIC THERAPY  
TRANSFUSION GUIDELINES  
USE OF HEMATOPOETIC GROWTH FACTORS  
WARFARIN

## **INFECTIOUS DISEASES**

ACALCULUS CHOLECYSTITIS  
ANTIBIOGRAMS  
BLOODSTREAM INFECTIONS  
CANDIDURIA  
CATHETER RELATED SEPTICEMIA  
CATHETER REPLACEMENT STRATEGIES  
ENDOCARDITIS PROPHYLAXIS  
ENDOCARDITIS DIAGNOSIS AND TREATMENT  
FEBRILE NEUTROPENIA  
FUO  
HIV+ PATIENT INFECTIONS  
MENINGITIS  
NECROTIZING SOFT TISSUE INFECTIONS  
NON-INFECTIOUS CAUSES OF FEVER  
OPHTHALMIC INFECTIONS  
PNEUMONIA, COMMUNITY ACQUIRED  
PNEUMONIA, HOSPITAL ACQUIRED  
SEPTIC SHOCK  
SINUSITIS  
SIRS  
TRANSPLANT INFECTION PROPHYLAXIS  
TRANSPLANT-RELATED INFECTIONS

## **NEUROLOGY**

AGITATION, ANXIETY, DEPRESSION & WITHDRAWAL



BRAIN DEATH  
GUILLAIN-BARRE SYNDROME  
INTRACEREBRAL HEMORRHAGE  
MYASTHENIA GRAVIS  
NEUROMUSCULAR COMPLICATIONS OF CRITICAL ILLNESS  
NON-TRAUMATIC COMA  
SEDATION  
STATUS EPILEPTICUS  
STROKE  
SUB-ARACHNOID HEMORRHAGE

### **PHARMACOLOGY**

AMINOGLYCOSIDE DOSING AND THERAPEUTIC MONITORING  
AMPHOTERICIN-B TREATMENT GUIDELINES  
ANALGESIA  
ANTIBIOTIC CLASSIFICATION & COSTS  
DRUG CHANGES WITH RENAL DYSFUNCTION  
PENICILLIN ALLERGY  
NEUROMUSCULAR BLOCKERS  
VANCOMYCIN  
THERAPEUTIC DRUG MONITORING

### **PULMONARY**

ARDS: HEMODYNAMIC MANAGEMENT  
ARDS: STEROID USE  
ARDS: VENTILATOR STRATEGIES  
ASTHMA  
BRONCHODILATOR USE IN VENTILATOR PATIENTS  
BRONCHOSCOPY & THORACENTESIS GUIDELINES  
COPD EXACERBATION & TREATMENT  
CXR (INDICATIONS)  
NONINVASIVE MODES OF VENTILATION  
ENDOTRACHEAL TUBES & TRACHEOTOMY  
TREATMENT OF AIRWAY OBSTRUCTION  
VENTILATOR WEANING PROTOCOL

## **RENAL**

ACUTE RENAL FAILURE :DIAGNOSIS  
ACUTE RENAL FAILURE :MANAGEMENT & TREATMENT  
DIALYSIS  
DIURETIC USE  
HYPERKALEMIA: ETIOLOGY & TREATMENT  
HYPERNATREMIA: ETIOLOGY & TREATMENT  
HYPOKALEMIA: ETIOLOGY & TREATMENT  
HYPONATREMIA: ETIOLOGY & TREATMENT  
OLIGURIA

## **SURGERY**

OBSTETRICAL COMPLICATIONS  
DISSECTING AORTIC ANEURYSM  
POST-OPERATIVE HYPERTENSION  
POST-OPERATIVE MYOCARDIAL ISCHEMIA (NON-CARDIAC  
ARRHYTHMIAS AFTER CARDIAC SURGERY  
POST-OPERATIVE BLEEDING  
POST-OPERATIVE MANAGEMENT OF ABDOMINAL  
POST-OPERATIVE MANAGEMENT OF OPEN HEART  
POST-OPERATIVE MANAGEMENT OF THORACOTOMY  
POST-OPERATIVE POWER WEANING  
POST-OPERATIVE MANAGEMENT OF CAROTID  
WOUND HEALING STRATEGIES

## **TOXICOLOGY**

ACETAMINOPHEN OVERDOSE  
ANAPHYLAXIS  
COCAINE TOXICITY  
ALCOHOL WITHDRAWAL  
HYPERTHERMIA  
LATEX ALLERGY  
UNKNOWN POISONING

## TRAUMA

ABDOMINAL COMPARTMENT SYNDROME  
BLUNT ABDOMINAL INJURY  
BLUNT AORTIC INJURY  
BLUNT CARDIAC INJURY  
DVT PROPHYLAXIS  
EXTREMITY COMPARTMENT SYNDROME  
HEAD INJURY  
HYPOTHERMIA  
IDENTIFICATION OF CERVICAL CORD INJURY  
SPINAL CORD INJURY  
OPEN FRACTURES  
PENETRATING ABDOMINAL INJURY  
PENETRATING CHEST INJURY

Referring to **Figure 21**, the acalculous cholecystitis decision support algorithm of the present invention is illustrated. If an intensivist suspects that acalculous cholecystitis may be present, the intensivist may not be certain of all of the aspects that would be indicative of this particular condition. Therefore, the intensivist is lead through a decision support algorithm, which first causes the intensivist to determine if the patient is clinically infected, either febrile or leukocytosis **800**. If this criteria is not met, the intensivist is prompted that it is unlikely that the patient has acalculous cholecystitis **802**.

If the patient is clinically infected **800**, the intensivist is prompted to determine whether the patient has had a previous cholecystectomy **804**. If patient has had a previous cholecystectomy, the intensivist is prompted that it is very unlikely that the patient has acalculous cholecystitis **806**. Alternatively, if a patient has not had a previous cholecystectomy, the intensivist is prompted to determine whether the patient has any of seven (7) risk factors, specifically: 1) Prolonged intensive care unit (ICU) stay (defined as greater than six (6) days); 2) recent surgery (particularly aortic cross clamp procedures); 3) hypotension; 4) positive end-

1 expiratory pressure (PEEP) greater than ten (10) centimeters (cm); 5) transfusion greater than six  
2 (6) units of blood; 6) inability to use the gastrointestinal (GI) tract for nutrition; or 7)  
3 immunosuppression (AIDS, transplantation, or leukemia) **808**. If the patient has none of these  
4 seven risk factors, the intensivist is prompted that the patient probably does not have acalculous  
5 cholecystitis **810**.

6 If the patient has any of the seven risk factors **808**, the intensivist is prompted to  
7 determine whether the patient has any of the following symptoms: right upper quadrant (RUQ)  
8 tenderness; elevated alkalinephosphatase; elevated bilirubin; or elevated liver transaminases  
9 **812**. If the patient has none of these four (4) symptoms **812**, the intensivist is prompted to  
10 consider other more likely sources of infection (see fever of unknown origin or FUO) **814**. If the  
11 infection remains undiagnosed following an alternative work-up, the intensivist is prompted to  
12 re-enter the algorithm **814**.

13 If the patient has any of these four (4) symptoms **812**, the intensivist is prompted to  
14 determine whether alternative intra-abdominal infectious sources are more likely **816**. If  
15 alternative intra-abdominal infectious sources are not more likely, the intensivist is prompted to  
16 determine whether the patient is sufficiently stable to go for a test **826**. If the patient is  
17 sufficiently stable to go for a test, the intensivist is prompted to perform an mso4  
18 Cholescintigraphy **836**. The normal AC is excluded **838**. If the test indicates an abnormality, the  
19 intensivist is prompted to consider a cholecystectomy or percutaneous drainage **840**. If the  
20 patient is not sufficiently stable to go for a test, the intensivist is prompted to perform a bedside  
21 ultrasound **828**. If no other infectious etiologies are identified and no abnormalities of the gall-  
22 bladder are noted but: a) the patient remains ill **830**, the intensivist is prompted to consider  
23 empiric cholecystostomy **832**. If no other infectious etiologies are identified and no  
24 abnormalities of the gall bladder are noted but: b) the patient is improving **830**, the intensivist is  
25 prompted to continue to observe the patient **834**.

26 If alternative intra-abdominal infectious sources are more likely **816**, the intensivist is

1 prompted to determine whether the patient is sufficiently stable to go for a test **818**. If the patient  
2 is sufficiently stable to go for a test **818**, the intensivist is prompted to perform an abdominal CT  
3 scan **820**. If no other infectious etiologies are apparent and the test: a) demonstrates  
4 abnormalities of the gall-bladder but not diagnostic; or b) no gall-bladder abnormalities are noted  
5 **822**, the intensivist is prompted to maintain continued observation of the patient **824**.  
6 Alternatively, if this criteria not met **822**, the intensivist is prompted to perform an mso4  
7 cholescintigraphy **836**. Normal AC is excluded **838**. If the test is abnormal, the intensivist is  
8 prompted to consider cholecystectomy or precutaneous drainage **840**. If the patient is not  
9 sufficiently stable to go for a test, the intensivist is prompted to perform a bedside ultrasound  
10 **828**. If no other infectious etiologies are identified and no abnormalities of the gall-bladder are  
11 noted but: a) the patient remains ill **830**, the intensivist is prompted to consider empiric  
12 cholecystostomy **832**. If no other infectious etiologies are identified and no abnormalities of the  
13 gall bladder are noted but: b) the patient is improving **830**, the intensivist is prompted to continue  
14 to observe the patient **834**.

15 Referring to **Figure 22**, the adrenal insufficiency decision support algorithm of the  
16 present invention is illustrated. When an intensivist suspects an adrenal problem may be  
17 presented in a patient, the intensivist may initiate the adrenal insufficiency decision support  
18 algorithm which prompts questions concerning all aspects of the condition. First the intensivist  
19 is prompted to determine whether the patient is either hypotensive and/or has been administered  
20 pressors for forty-eight hours or longer **900**. If neither condition is met, the system advises the  
21 intensivist that it is unlikely that an adrenal problem is present **902**.

22 If one or both conditions are met, the intensivist is asked whether an obvious cause for  
23 hypotensive blood pressure or treatment with pressors are manifested, such as hypovolemia or  
24 low blood volume, myocardial dysfunction, or spinal injury **904**. If at least one of these obvious  
25

causes is present, the intensivist is alerted by the system that the underlying cause must first be treated **906**. If treatment of a suspected underlying cause is reversed, yet the hypotension or pressor need persists, the intensivist is further directed to determine whether other adrenal problems have occurred in the patient's history **908, 910, 912**

In order to examine prior treatment issues, the intensivist is first prompted by the system to determine if the patient has been treated with steroids in the previous six months for at least a two week period **908**. Next, the intensivist is prompted to determine whether the patient has hyponatremia or hyperkalemia **910**. The intensivist is also prompted to determine whether the patient has experienced anticoagulation or become coagulopathic prior to the hypotension or pressor treatment **912**. According to the responses provided by the intensivist to the system queries or blocks **908, 910, and 912**, the system calculates a treatment action **914** as follows: The array of possible responses to diagnosis questions **908, 910, and 912** are given a Decision Code as shown in Table 1: Adrenal Insufficiency Considerations, below.

Table 1: Adrenal Insufficiency Considerations

Question 1	Question 2	Question 3	Decision Code
908	910	912	
N	N	N	A
N	N	Y	A
N	Y	N	B
N	Y	Y	C
Y	Y	Y	C
Y	N	N	D
Y	Y	N	B

Y	N	Y	D
Y	Y	Y	C

1

2 The possible decision codes of Table 1 are as follows:

Decision Code	Treatment Action
A	Do cosyntropin stim test
B	Consider possible Adrenal Insufficiency. Give decadron 5 mg IV, so cosyntropin stim test and empirically treat with hydrocortione 50 mg IV every 8 hours until stim test results return.
C	Consider possible Adrenal Insufficiency, secondary to adrenal hemorrhage. Give decadron 5 mg IV, so cosyntropin stim test and empirically treat with hydrocortione 50 mg IV every 8 hours until stim test results return.
D	Do cosyntropin stim test, may empirically treat with hydrocortisone 25-50 mg IV every 8 hours until stim test results return

3

4 Besides specialized treatment actions listed in the decision codes above, the intensivist is  
5 directed to administer a cosyntropin stimulation test **914** in order to see how much cortisone the  
6 adrenal gland is producing.

7 After performing the cosyntropin stimulation test, the intensivist is prompted to enter the  
8 patient's level of cortisol before administering cosyntropin and thirty minutes afterwards **916**.

9 The software analyzes the test results as follows:

10 The results in Table 2, shown below, are shown as having certain decision codes A through F.

11 **Table 2: Cosyntropin Stimulation Test Results**

<b>basal (A)</b>  <b>&lt; 15</b>	basal (B) 15-20	basal (C) > 25
stim (D) < 5	stim (E) 5-10	stim (F) > 10

12

Depending upon the outcome of the analysis of Table 2, one of the treatment actions, shown below in Table 3, will be displayed 918.

**Table 3: Cosyntropin Test Result Treatment Actions**

Decision Code	Treatment Action
A + D	<u>Adrenal insufficiency diagnosed - treat with hydrocortisone 50 mg IV every 8 hours and consider endocrine consult</u>
A + E B + D	Probable Adrenal insufficiency- treat with hydrocortisone 25-50 mg IV every 8 hours and taper as intercurrent illness improves
A + F B + E	Possible Adrenal insufficiency- consider treatment with hydrocortisone 25 mg IV every 8 hours and taper as intercurrent illness improves
A + F B + F C + E C + F	Adrenal insufficiency unlikely- would not treat

Referring to **Figure 23**, the blunt cardiac injury decision support algorithm of the present invention is illustrated. If an intensivist suspects that blunt cardiac injury may be present, the intensivist may not be certain of all aspects that would be critical to or indicative of this particular condition. Therefore, the intensivist is lead through a decision support algorithm, which first causes the intensivist to determine whether any of seven (7) risk factors are present:

1) was thoracic impact greater than fifteen (15) mph; 2) was the steering wheel deformed; 3) was there precordial ecchymosis, contusions, or abrasions; 4) was marked precordial tenderness present; 5) was there a fractured sternum; 6) were bilateral rib/costal cartilage fractures present; 7) were thoracic spine fractures present 1000. If none of the 7 risk factors are present, the intensivist is prompted that no further evaluation is necessary 1002. If any of the 7 risk factors are present, the intensivist is prompted to obtain an electrocardiogram (ECG) and chest X-ray (CXR) 1004.



Once the results of the ECG and CXR are obtained, the intensivist is prompted to determine: whether the ECG results are abnormal, with abnormal being defined as anything other than sinus rhythm, including ectopy and unexplained sinus tachycardia (greater than 100 beats/minute); and whether the CXR results are abnormal, with abnormal being defined as any skeletal or pulmonary injury, especially cardiac enlargement **1006**. If either the ECG or CXR are not abnormal, the intensivist is prompted that a monitored bed is unnecessary for the patient **1008**. If either the ECG or CXR are abnormal, the intensivist is prompted to determine whether there is any hemodynamic instability (hemodynamic instability being defined as the absence of hypovolemia, spinal cord injury, or sepsis) that cannot be explained by hypovolemia, spinal cord injury, or sepsis **1010**.

If this criteria is not met, the intensivist is prompted: that the patient should be in a monitored bed; that the ECG should be repeated at 24 hours; that, at any time, if unexplained hemodynamic instability is present, the intensivist should request a stat echo; and that, if blunt thoracic aortic injury is also suspected, a transesophageal echocardiogram (TEE) is favored over a transthoracic echocardiogram (TTE) **1012**. Once the results of these tests are obtained, the intensivist is prompted further to determine whether ectopy, arrhythmia, or abnormality is present on the ECG **1014**. If none of these criteria are met, the intensivist is prompted that cardiac injury is excluded **1016**. If any of these criteria are met, the intensivist is prompted that he should consider monitoring the patient for an additional 24 hours **1018**.

If the internist determines that there is any hemodynamic instability that cannot be explained by hypovolemia, spinal cord injury, or sepsis **1010**, he is prompted: to perform a stat echo; and, if blunt thoracic aortic injury is also suspected, that a transesophageal echocardiogram (TEE) is favored over a transthoracic echocardiogram (TTE) **1020**. Once the results of the stat

1 echo are obtained, the intensivist is prompted to determine whether the echo is abnormal with  
2 possible causes for the abnormality being: pericardial effusion (tamponade; hypokineses or  
3 akinesis (wall motion); dilatation or reduced systolic function; acute valvular dysfunction; and/or  
4 chamber rupture **1022**. If the stat echo is abnormal, the intensivist is prompted to treat as  
5 indicated for the particular cause of the abnormality **1026**. If the stat echo is not abnormal, the  
6 intensivist is prompted to continue to monitor the patient and repeat the ECG at 24 hours **1024**.

1           Once the results of the ECG are obtained, the intensivist is prompted to determine  
2 whether ectopy, arrhythmia, or abnormality are present on the ECG **1014**. If this criteria is not  
3 met, the intensivist is prompted that cardiac injury is excluded **1016**. If this criteria is met, the  
4 intensivist is prompted that he should consider monitoring the patient for an additional 24 hours  
5 **1018**.

6           Referring to **Figure 24**, the candiduria decision support algorithm, which is yet another  
7 decision support algorithm of the present invention is illustrated. In the candiduria decision  
8 support algorithm, the intensivist is presented with the criteria for diagnosing candiduria, or  
9 severe fungal infection. First, the intensivist determines whether the patient has any medical  
10 conditions that render the patient prone to fungal infections, such as diabetes, GU anatomic  
11 abnormality, renal transplant, or pyuria **1100**. If there are no such conditions, the intensivist is  
12 next prompted by the system to look for dissemination or spreading of the fungal infection **1102**.  
13 If the infection does not seem to have spread, the intensivist is prompted to change the patient's  
14 catheter and test for pyuria after twenty four hours have passed **1104**.

15           The intensivist is prompted by the system to determine whether the patient can have P.O.  
16 **1106**. If the patient can take P.O., the system next prompts the intensivist to determine whether  
17 azoles, an organic compound for inhibiting fungal growth, have been administered in the past  
18 three days to fight the infection **1108**. If azoles have been previously administered, the systemic  
19 infection diagnosis is confirmed and the intensivist is referred to the systemic amphotericin  
20 dosing algorithm **1110**. If azoles have not been previously administered, directions for the  
21 proper treatment dosage of fluconazole (a type of azole) is provided to the intensivist along with  
22 adjustments for the species of fungus found **1112**. Where the patient cannot take P.O., the

1 intensivist is again referred to the systemic amphotericin dosing algorithm **1114**.

2 When the patient does have some condition prone to fungal infection, the intensivist is  
3 prompted to determine what other signs of dissemination are exhibited in the patient **1116**. The  
4 intensivist is prompted to see if the patient can take P.O. If the patient cannot take P.O., the  
5 intensivist is referred to the systemic amphotericin dosing algorithm **1120**. If the patient can take  
6 P.O., the intensivist is prompted to check whether azoles have been administered in the previous  
7 three days **1122**. If azoles have been administered, the systemic infection is confirmed and the  
8 intensivist is referred to the systemic amphotericin dosing algorithm **1124**. If no azoles have  
9 been administered previously, the intensivist is given instructions for administering fluconazole  
10 to treat the fungal infection **1126**.

11 If there is no evidence of dissemination, the intensivist is still prompted to determine  
12 whether the patient can take P.O. **1128**. Where the patient cannot take P.O., directions are  
13 provided to administer amphotericin bladder washing procedures **1130**. If the patient cannot take  
14 P.O., the intensivist is prompted to determine whether azoles have been administered in the  
15 previous three days **1132**. If azoles have been administered, the systemic infection is confirmed  
16 and the intensivist is referred to the systemic amphotericin dosing algorithm **1134**. If no azoles  
17 have been administered previously, the intensivist is given instructions for administering  
18 fluconazole to treat the fungal infection **1136**.

19  
20 Referring to **Figure 25**, the Cervical Spine Injury decision support algorithm of the  
21 present invention is illustrated. If an intensivist suspects that a cervical spine injury may be  
22 present, the intensivist may not be certain of all of the factors that would be indicative of this

particular condition. Therefore, the intensivist is lead through a decision support algorithm, which first prompts the intensivist to determine if the patient is awake, alert, not intoxicated, and has no mental status changes **1200**. If this criteria is met, the intensivist is prompted to determine whether the patient has any neck pain **1202**. If the patient does not have any neck pain, the intensivist is prompted to determine whether the patient has any other pain which would distract from their neck pain **1204**. If this criteria is not met, the intensivist is prompted to determine whether the patient has any neurologic deficits **1206**. If this criteria is not met, the intensivist is prompted that a stable C-spine is present if the patient can flex, extend, move neck left/right without pain and without neck tenderness to palpitation **1208**. The intensivist is prompted further that he can remove the collar **1208**.

Alternatively, if the patient does have neck pain **1202**, the intensivist is prompted to order 3 x rays **1210** consisting of: 1) lateral view revealing the base of the occiput to the upper border of the first thoracic vertebra; 2) anteroposterior view revealing spinous processes of the second cervical through the first thoracic vertebra; and 3) an open mouth odontoid view revealing the lateral masses of the first cervical vertebra and entire odontoid process **1210**. If the x rays are normal the intensivist is prompted to consider extension then flexion lateral x rays; if normal he is prompted that he can remove the collar; if abnormal, he is prompted to obtain a surgical consult **1212**. If the x rays are abnormal, the intensivist is prompted to obtain a surgical consult and order a CT scan **1214**. If the x rays are indeterminate, the intensivist is prompted to order a CT scan **1216**.

Alternatively, if the patient has no other pain which would distract from their neck pain **1204**, the intensivist is prompted to order 3 x rays (the same types of x rays described in **1210** above with the same prompting based on normal, abnormal, or indeterminate x rays) **1218**.

If the patient does have neurologic deficits **1206**, the intensivist is prompted to determine whether the neurologic deficit is referable to the cervical spine **1226**. If this criteria is not met, the intensivist is prompted to order 3 x rays (the same types of x rays described in **1210** above with the same prompting based on normal, abnormal, or indeterminate x rays) **1218**. If the neurologic deficit is referable to the cervical spine **1226**, the intensivist is prompted that the patient should obtain immediate spine trauma surgery consult and CT or MRI (if available) **1228**.

Alternatively, if the intensivist determines that the patient does not pass the criteria of being awake, alert, not intoxicated and having no mental status changes **1200**, the intensivist is prompted to determine whether the patient has severe head trauma **1232**. If this criteria is met, the intensivist is prompted to order CT of the neck with head CT **1236**. If this criteria is not met, the intensivist is prompted to determine whether the patient has any neurologic deficit referable to the cervical spine **1234**. If the intensivist determines that the patient does have a neurologic deficit referable to the cervical spine, the intensivist is prompted that the patient should obtain immediate spine trauma surgery consult and CT or MRI (if available) **1228**. If the intensivist determines that the patient does not have a neurologic deficit referable to the cervical spine **1234**, he is prompted to order 3 x rays (the same types of x rays described in **1210** above with the same prompting based on normal, abnormal, or indeterminate x rays) **1218**.

Referring to **Figure 26**, the Oliguria decision support algorithm of the present invention is illustrated. If an intensivist suspects that Oliguria may be present, the intensivist may not be certain of all of the aspects that would be indicative of this particular condition. Therefore, the intensivist is lead through a decision support algorithm, which first causes the intensivist to determine if the patient is oliguric, with the criteria being passage of less than 25 cc of urine in a period of 2 hours **1300**. If this criteria is met the intensivist is prompted to determine whether the patient is anuric (the criteria for which is passage of less than 10 cc of urine in a 2 hour period) in spite of fluid administration **1302**.

If this criteria is met, the intensivist is prompted to determine whether the urinary catheter is working by flushing the catheter **1304**. The intensivist is then prompted to determine whether the catheter is functioning **1306**. If the catheter is not functioning, the intensivist is prompted to replace or reposition the catheter **1308**. If the catheter is functioning, the intensivist is prompted to determine whether the patient has a history of: 1) renal stone disease; 2) abdominal, pelvic, or retroperitoneal cancer; or 3) recent pelvic or retroperitoneal surgery **1310**. If any of these criteria are met, the intensivist is prompted to perform the following actions: 1) do renal ultrasound emergently to rule out obstruction; 2) while waiting for ultrasound, administer fluid at the rate of 7-15 ml/kg of bodyweight; and 3) send urine for specific gravity determination **1312**. Based on the renal ultrasound test results, the intensivist is prompted to determine whether an obstruction is present **1314**. If an obstruction is determined to be present, the intensivist is prompted to consult a urologist immediately **1316**.

Alternatively, if the intensivist determines that the patient does not have a history of: 1) renal stone disease; 2) abdominal, pelvic, or retroperitoneal cancer; or 3) recent pelvic or

retroperitoneal surgery **1310**, the intensivist is prompted to determine whether: 1) the patient has a history of heart failure or known ejection fraction of less than 30 percent; or 2) there are rales on the physical exam **1318**.

Alternatively, if following the renal ultrasound test, the intensivist determines that there is no obstruction the intensivist is prompted to determine whether: 1) the patient has a history of heart failure or known ejection fraction of less than 30 percent; or 2) there are rales on the physical exam **1318**.

If the intensivist determines that the patient is not anuric **1302**, then the intensivist is prompted to determine whether: 1) the patient has a history of heart failure or known ejection fraction of less than 30 percent; or 2) whether there are rales on the physical examination **1318**. If this criteria is not met, the intensivist is prompted to administer fluids to the patient at the rate of 10-20 ml/kg of bodyweight **1320** and send the patient's urine sample for a specific gravity test **1322** as more fully described in **Figure 26A**.

Alternatively, if the patient does: 1) have a history of heart failure or known ejection fraction less than 30 percent; or 2) there are rales on the physical exam **1318**, the intensivist is prompted to determine whether there has been a chest x-ray (CXR) in the last 6 hours **1324**. If this criteria is not met, the intensivist is prompted to determine whether there has been a change in respiratory status **1326**. If there has been no change in the respiratory status, the intensivist is prompted to administer 7-15 ml of fluids per kg of bodyweight **1328** and to send the patient's urine sample for a specific gravity test.

Alternatively, if the intensivist determines that there has been a change in respiratory status **1326**, the intensivist is prompted to: 1) do a chest x-ray; and 2) determine whether there is



1 evidence of edema or congestion **1334**. If there is evidence of edema or congestion **1334**, the  
2 intensivist is prompted to: 1) insert a PA catheter to measure wedge pressure and liver function  
3 to direct fluid replacement; and 2) send urine creatinine and sodium **1332**.

4 If the intensivist determines that there has been a CXR in the last 6 hours **1324**, the  
5 intensivist is prompted to determine whether there is evidence of edema or congestion **1330**. If  
6 there is no evidence of edema or congestion, the intensivist is prompted to administer 7-15 ml of  
7 fluids per kg of bodyweight **1328** and send the patient's urine for a specific gravity test **1322**.

8 Alternatively, if the intensivist determines there is evidence of edema or congestion **1330**,  
9 the intensivist is prompted to: : 1) insert a PA catheter to measure wedge pressure and liver  
10 function to direct fluid replacement; and 2) send urine creatinine and sodium **1332**.

11 Referring now to **Figure 26A**, the oliguria algorithm description continues. Following  
12 the specific gravity test of the patient's urine, the intensivist is prompted to determine whether  
13 the results indicate the specific gravity is less than 1.018. If this criteria is met, the intensivist is  
14 prompted to: 1) send blood and urine immediately to test for blood urea nitrogen (BUN),  
15 creatinine, electrolytes, and Hgb, and spot urine for creatinine, sodium, and sediment; and 2)  
16 administer 5-10 ml of fluid per kg of bodyweight **1356**. Once the results of these tests are  
17 obtained, the intensivist is prompted to determine what is the Hgb **1338**.

18 If the Hgb has increased by more than 1.5 gm/dl compared to the previous hgb **1340**, the  
19 intensivist is prompted to: 1) administer fluids 5-10 ml/kg of bodyweight and follow the urine  
20 output closely **1342**. Following this, the intensivist is prompted to determine whether the labs  
21 confirm renal failure by use of the formula  $FE_{Na} = \frac{\text{Urine Na} \times \text{Serum Creatinine}}{\text{Urine Creatinine} \times \text{Serum Na}} \times 100$  **1344**.

1 If the Hgb is within 1.5 gm/dl from the previous hgb or no comparison **1352**, the  
2 intensivist is prompted to determine what is the mean blood pressure **1354**. If the mean blood  
3 pressure is determined to be within 20 percent or higher than the baseline blood pressure **1356**,  
4 the intensivist is prompted to determine whether the labs confirm renal failure **1344**. If the mean  
5 blood pressure is determined to be greater than 20 percent below the baseline pressure **1358**, the  
6 intensivist is prompted to give additional fluids and consider invasive hemodynamic monitoring  
7 **1360**. Following this, the intensivist is prompted to determine whether the labs confirm renal  
8 failure by use of the formula  $FE_{Na} = \text{Urine Na} \times \text{Serum Creatinine} / \text{Urine Creatinine} \times \text{Serum Na} \times$   
9 100 **1344**.

10 Alternatively if the Hgb has decreased by 1.5 gm/dl compared to the previous hgb **1362**,  
11 the intensivist is prompted to: 1) transfuse PRBCs as needed; 2) look for source of bleeding and  
12 check PT, aPTT, & platelet count **1364**. Following this, the intensivist is prompted to determine  
13 what is the mean blood pressure **1354**. If the mean blood pressure is determined to be greater  
14 than 20 percent below the baseline pressure **1358**, the intensivist is prompted to give additional  
15 fluids and consider invasive hemodynamic monitoring **1360**. Following this, the intensivist is  
16 prompted to determine whether the labs confirm renal failure by use of the formula  $FE_{Na} = \text{Urine}$   
17  $\text{Na} \times \text{Serum Creatinine} / \text{Urine Creatinine} \times \text{Serum Na} \times 100$  **1344**.

18 If the labs do not confirm renal failure, as indicated by  $FE_{Na} \leq 1$  percent **1346**, the  
19 intensivist is prompted to: 1) continue to administer fluids and follow urine output; and 2)  
20 recheck creatinine in 6-12 hours **1348**.

21 Alternatively, if the labs do confirm renal failure, as indicated by  $FE_{Na} > 1$  percent **1350**,  
22 the intensivist is prompted to: 1) place central venous pressure (CVP); 2) Assure adequate

1 intravascular volume; 3) give trial of diuretics: 40 mg lasix IV, if no response in 1 hour, give  
2 hydrodiuril 500 mg IV, wait 20-30 minutes then give 100 mg lasix, if persistent oliguria, restrict:  
3 1) fluids; 2) potassium & phosphate; if diuresis ensues, restrict only potassium & phosphate; in  
4 both situations, adjust all renally excreted medications; and 4) see acute renal failure **1350**.

5 Referring now to **Figure 26B**, the oliguria algorithm description continues.  
6 Alternatively, following the specific gravity test of the patient's urine, the intensivist is prompted  
7 to determine whether the results indicate the specific gravity is greater than or equal to 1.018  
8 **1336**. If this criteria is not met **1364**, the intensivist is prompted to determine whether the urine  
9 is dark or tea colored **1366**. If this criteria is met, the intensivist is prompted to: 1) check  
10 creatinine phospho/kinase; and 2) force fluids to induce diuresis **1368**.

11 If the intensivist determines that the urine is not dark or tea colored, the intensivist is  
12 prompted to: 1) administer 10-20 ml of fluids per kg of bodyweight; and 2) check hgb **1370**. The  
13 intensivist is then prompted to determine what is the hgb **1372**.

14 If the hgb is determined to be greater than 1.5 gm/dl higher than the previous hgb **1374**,  
15 the intensivist is directed to: 1) force fluids; and 2) continue to follow the urine output **1376**.

16 Alternatively, if the hgb is determined to be within 1.5 gm/dl of the last hgb or there is no  
17 hgb for comparison **1378**, the intensivist is prompted to determine what is the mean blood  
18 pressure **1380**. If the mean blood pressure is determined to be 20 percent or higher than the  
19 baseline pressure **1382**, the intensivist is prompted to: 1) continue to administer fluids; 2) follow  
20 urine output; and 3) check creatinine in 6-12 hours **1384**. If the mean blood pressure is  
21 determined to be greater than 20 percent below the baseline pressure **1386**, the intensivist is

1 prompted to: 1) continue to push fluids; 2) consider invasive hemodynamic monitoring; and 3) if  
2 post-op abdominal trauma, consider abdominal compartment syndrome **1388**.

3 If the hgb is determined to be greater than 1.5 gm/dl below the previous hgb **1390**, the  
4 intensivist is prompted to: 1) transfuse blood as needed; 2) look for bleeding source; 3) check  
5 PT, aPPT & platelet count; 4) continue to push fluids; and 5) recheck hgb in 1-2 hours **1392**.

6 Referring to **Figure 27**, the open fractures decision support algorithm of the present  
7 invention is illustrated. Open fractures are where bone, cartilage, or a tooth break and push  
8 through the skin surface. The intensivist is first prompted by the system to determine whether  
9 the patient has an open fracture **1500**. If one has occurred, the intensivist must then determine  
10 whether the wound is contaminated with soil, or was inflicted in a barnyard **1502** in order to  
11 address higher risk of infection. If the wound is contaminated with soil, or was inflicted in a  
12 barnyard, the intensivist is prompted to administer a high dose of penicillin to the antibiotics  
13 prescribed **1504**. The intensivist is also prompted to take several treatment steps **1506**. These  
14 treatment steps include administering tetanus prophylaxis, such an antitoxin injection,  
15 monitoring staphylococcus aureus until twenty-four hours after surgery, caring for the wound  
16 within six hours, and where the injury is found to be more severe during surgery, the intensivist  
17 is prompted to administer aminoglycosides for seventy two hours.

18 If the wound is not contaminated with soil, or was inflicted in a barnyard, the intensivist  
19 is next prompted to determine the severity of the wound **1508**. To do so, the intensivist must  
20 determine the length of the wound and corresponding soft tissue damage. If the wound is either  
21 less than one centimeter and clean or greater than a centimeter long without extensive soft tissue  
22 damage, the Intensivist is prompted to take several treatment steps **1506** as previously described.

1 Where the soft tissue damage is extensive or amputation has occurred, the intensivist is  
2 prompted by the system to make further determinations **1510**, **1512**, **1514** about the wound  
3 caused by the fracture. The intensivist is prompted to determine if enough soft tissue coverage is  
4 remaining for the wound to close and heal **1510**, if any arterial repair is needed **1512**, and if  
5 extensive soft tissue damage with periosteal injury, and bone exposure **1514**. If there is  
6 adequate soft tissue coverage, the intensivist is advised that risk of infection is low and directed  
7 to take treatment actions **1516**. If arterial damage requiring repair is present, the intensivist is  
8 advised by the system that risk of infection is moderate to high and given treatment instructions  
9 **1518**. Where there is soft tissue injury with periosteal stripping and bone exposure, the  
10 intensivist is alerted by the system that risk of infection is high and given treatment instructions  
11 **1520**. The treatment instructions in each case **1516**, **1518**, **1520** include administering tetanus  
12 prophylaxis, such as an antitoxin injection, caring for the wound within six hours, and performing:  
13 monitoring for staphylococcus aureus, and administering aminoglycosides and high doses of  
14 penicillin, all for seventy two hours before and after any operative procedures.

15 If the intensivist has determined that no exposed fracture has occurred, the system next  
16 prompts the intensivist to determine whether there is any evidence of neuro-vascular damage  
17 **1522**. If there is evidence of neuro-vascular damage, the intensivist is prompted to consult with a  
18 neurosurgeon or vascular surgeon immediately **1524**. If the intensivist determines there is no  
19 evidence of neuro-vascular damage to the patient, the system next prompts the intensivist to  
20 determine whether the patient has compartment syndrome **1526**. If there is evidence of  
21 compartment syndrome seen in the patient, the intensivist is prompted to consult orthopedics  
22 right away **1528**. If there is no evidence of compartment syndrome seen in the patient, the

intensivist is still prompted to consult orthopedics, but without any prompt for time sensitivity  
1530.

Referring to **Figure 28**, the Pancreatitis diagnostic algorithm of the present invention is  
illustrated. To evaluate whether a patient has pancreatitis, the intensivist is first prompted to  
examine whether severe epigastric abdominal pains and amylase levels three times greater than  
normal are present in the patient **1600**. If neither or one of the conditions is present, the  
intensivist is prompted to consider other causes of the abdominal pain, such as mesenteric  
ischemia, a perforated ulcer, intestinal obstruction, biliary colic, or an ectopic pregnancy **1602**.

If severe epigastric abdominal pains and amylase levels three times greater than normal  
are present, the intensivist is next prompted to provide the Ranson Criteria which is a criteria  
associated with the severity of pancreatitis and the potential outcome or prognosis at that  
particular level of severity, or Apache II score which is also a score associated with the severity  
of the disease and the potential prognosis at a particular level of the patient **1604**. If the patient  
has a Ranson Criteria less than three or an Apache II score of less than eight, the intensivist is  
prompted by the system to consider removing the patient from the Intensive Care Unit **1606**.  
However, if the patient has a Ranson Criteria greater than three or an Apache II score of greater  
than eight, the intensivist is instructed to perform an abdominal ultrasound test within twenty-  
four hours **1607**. If the results of the ultrasound test show a biliary obstruction, the intensivist is  
instructed to consider performing an ERCP to find and remove any gallstones **1608**.

If the abdominal ultrasound results do not show any biliary obstruction, intensivist is next  
prompted to perform more diagnostic tests **1610**. The intensivist is directed to perform a  
Dynamic IV contrast and an abdominal Computerized Tomography (CT) scan. If the intensivist

1 does not suspect a surgical condition exists, such as a perforated ulcer, mesenteric infarction or  
2 pancreatic infection, the tests may be performed after three days have passed. If the intensivist  
3 does suspect a surgical condition exists, the tests should be performed within three days. In  
4 either case, if the patient has creatinine levels greater than or equal to 2 milligrams per dl, the  
5 intensivist should not perform the Dynamic IV contrast test.

6 Once the CT scan is performed, the intensivist is prompted to determine whether  
7 necrotizing pancreatitis is present **1612**. The intensivist is next required to determine whether  
8 the patient has improved since admission **1614**. If no improvement has been seen, the intensivist  
9 is directed to perform percutaneous fluid aspiration and do a gram stain culture the collected  
10 fluid **1616**. If the culture shows infection **1618**, the intensivist is directed to perform surgical  
11 debridement of the pancreas **1620**. If the results of the culture are sterile **1622**, the intensivist is  
12 directed to closely follow up on the patient's condition **1624** and watch for clinical deterioration  
13 **1626**. If the patient does further deteriorate, the intensivist is then instructed to perform a  
14 surgical debridement of the pancreas **1628**. If the patient does not deteriorate, the intensivist is  
15 still prompted to closely follow the patient's condition **1630**.

16 Where the CT scan does not show signs of necrotizing pancreatitis **1612**, the intensivist is  
17 prompted by the system to closely observe the patient **1632**. The intensivist is also prompted to  
18 check whether clinical deterioration is occurring **1634**. If no deterioration is observed, the  
19 intensivist continues to observe the patient's condition **1636**. If clinical deterioration is occurring  
20 **1634**, the intensivist is directed to perform percutaneous fluid aspiration and do a gram stain  
21 culture the collected fluid **1616**. If the culture shows infection **1618**, the intensivist is directed to  
22 order surgical debridement of the pancreas **1620**. If the results of the culture are sterile **1622**, the

1 intensivist is directed to closely follow up on the patient's condition **1624** and watch for clinical  
2 deterioration **1626**. If the patient does further deteriorate, the intensivist is then prompted to  
3 order a surgical debridement of the pancreas **1628**. If the patient does not deteriorate, the  
4 intensivist is still directed by the system to closely follow the patient's condition **1630**.

5 Referring to **Figure 29**, the penicillin allergy diagnosis algorithm of the present invention  
6 is illustrated. In order to diagnose a penicillin allergy, the intensivist is first prompted to  
7 determine whether the patient has a history suggestive of previous penicillin or cephalosporin  
8 anaphylaxis **1700**. Various known reactions, including angioedema, flushing, pruritis, airway  
9 obstruction, syncope, and hypertension, are displayed for the intensivist's review. If the patient  
10 has previously had any of these reactions, the intensivist is prompted to determine whether the  
11 patient has ever taken synthetic or partially synthetic antibiotics, such as ampicillin, amoxicillin,  
12 duricef or kefzol, without any anaphylaxis symptoms **1702**. If the patient has taken synthetics  
13 without reaction, the intensivist is advised by the system that penicillin or cephalosporin may be  
14 administered **1716**. If the patient has reacted to synthetic or partially synthetic antibiotics, the  
15 intensivist is next prompted to determine whether the patient needs penicillin or cephalosporin  
16 specifically **1704**.

17 If the patient is not required to have penicillin or cephalosporin, the intensivist is  
18 prompted to administer the synthetic antibiotics **1706**. If the patient does need penicillin or  
19 cephalosporin, the intensivist is directed by the system to consider consulting with an allergist or  
20 immunologist and perform skin tests for reactions **1708**. Next, the intensivist is prompted to  
21 enter whether the skin test was positive **1710**. If the results are negative, the intensivist is further  
22 directed by the system to administer penicillin or cephalosporin with caution, to consider



pretreatment with benadryl or prednisone to counter any reaction, and to closely monitor the patient **1712**. If the results of the skin test are positive, the intensivist is prompted by the system to perform desensitization procedures **1714**.

If the patient does not have a history suggestive of previous penicillin or cephalosporin anaphylaxis **1700**, the intensivist is prompted to determine whether the patient has previously experienced skin-level reactions, such as exfoliative dermatitis, Stevens Johnson Syndrome, or Toxic Epidermal Necrolysis, when given penicillin or cephalosporin **1718**. If the patient has previously experienced one of these reactions, the intensivist is directed by the system to administer an alternative antibiotic **1720**. If the patient has not experienced one of these reactions, the intensivist is prompted to determine whether there is a history of any rash when given penicillin or cephalosporin **1722**. If the patient has not previously had a rash when given penicillin or cephalosporin, the intensivist is advised that the patient will most likely be able to take penicillin or cephalosporin **1724**.

If the patient has previously experienced a rash when given penicillin or cephalosporin, the intensivist is prompted to determine whether the rash presented when the patient was given ampicillin or amoxycillin **1726**. If the rash resulted from ampicillin or amoxycillin, the intensivist is next prompted to determine whether the rash was urticarial **1728**. If the rash was not urticarial, the intensivist is advised by the system that the patient probably can take penicillin or cephalosporin, but should be closely monitored **1730**. If the rash was urticarial, the intensivist is prompted to determine whether or not the patient needs penicillin or cephalosporin **1704**.

If the patient is not required to have penicillin or cephalosporin, the intensivist is directed by the system to administer the synthetic antibiotics **1706**. If the patient does need penicillin or

1 cephalosporin, the intensivist is directed to consider consulting with an allergist or immunologist  
2 and perform skin tests for reactions **1708**. Next, the intensivist is prompted to enter whether the  
3 skin test was positive **1710**. If the results are negative, the intensivist is further directed to  
4 administer penicillin or cephalosporin with caution, to consider pretreatment with benadryl or  
5 prednisone to counter any reaction, and to closely monitor the patient **1712**. If the results of the  
6 skin test are positive, the intensivist is directed to perform desensitization procedures **1714**.

7 Referring to **Figure 30**, the Post-Op Hypertension decision support algorithm of the  
8 present invention is illustrated. If an intensivist determines that there may be a possibility of  
9 post-op hypertension, the intensivist may not be certain of all aspects that would be involved in  
10 this particular condition. Therefore, the intensivist is lead through a decision support algorithm  
11 which prompts the intensivist to determine the appropriate care to be given.

12 Initially, the intensivist is prompted to determine whether the patient is hypertensive (BP  
13 greater than 20 percent above mean baseline) **1800**. If this criteria is met, the intensivist is  
14 prompted to determine whether the patient has any of the causes of reversible hypertension: 1)  
15 hypercapnia; 2) bladder distension; 3) pain; 4) increased ICP; 5) drugs (pressors, cocaine,  
16 ketamine and chronic MAO use with indirect acting vasopressors); 6) automatic hyperreflexia; or  
17 7) volume overload **1802**. If any of these criteria are met, the intensivist is prompted to first treat  
18 those specific etiologies and, if pressure remains high, re-enter algorithm **1804**.

19 Alternatively, if none of these criteria are met **1802**, the intensivist is prompted to  
20 determine whether the patient is at risk of injury from post-op hypertension (i.e., vascular  
21 surgery, coronary artery disease, neurosurgery, ocular surgery, etc.) **1806**. If this criteria is not  
22 met **1806**, the intensivist is prompted to determine whether the BP is greater than 40 percent

1 above mean baseline **1808**. If this criteria is not met, the intensivist is prompted that the patient  
2 may not need BP treatment **1810**.

3 If the BP is greater than 40 percent above the mean baseline **1808**, the intensivist is  
4 prompted to determine whether the patient is in pain **1812**. If this criteria is met **1812**, the  
5 intensivist is prompted to treat pain and continue **1814**. Following this prompt **1814**, the  
6 intensivist is prompted next to determine whether the patient is actively bleeding or at significant  
7 risk for post-op bleeding (i.e., “moist closure” or high drain output) **1816**. If this criteria is met  
8 **1816**, the intensivist is prompted to use only short acting agents including emolol and  
9 nitroprusside as needed until bleeding has abated **1818**.

10 Alternatively, if this criteria is not met **1816**, the intensivist is prompted to determine  
11 whether the patient is tachycardic (absolute greater than 90 bpm or ((relative greater than 15  
12 percent over baseline)) **1820**. If this criteria is met **1820**, the intensivist is prompted to go to  
13 Decision Table C, which is programmed for the condition of a high heart rate. If this criteria is  
14 not met **1820**, the intensivist is prompted to eliminate (NOT C) Table C and proceed to the next  
15 decision point **1820**.

HR↑Table C

	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 <sup>ST</sup>	L	E	L	L	A	E
	2 <sup>ND</sup>	E	L	A	N	N	A

The intensivist is prompted next to determine whether the patient is bradycardic (absolute less than 60 bpm) **1822**. If this criteria is met, the intensivist is prompted to go to Decision Table B, which is programmed for the condition of a low heart rate.

HR ↓ Table B

	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 <sup>ST</sup>	N	N	A	N	A	A
	2 <sup>ND</sup>	S	S	S	H	H	H

If this criteria is not met, the intensivist is prompted to eliminate (NOT B) Table B and proceed to the next decision point **1822**. [Note: If NOT C and NOT B, the intensivist is prompted to go to Table A by default, i.e., If NOT C and NOT B Then A].

HR (nl) Table A

	CAD	Y	Y	Y	N	N	N
	RAD	N	Y	Y	N	Y	N
	↓EF	N	N	Y	N	Y	Y
Treatment	1 <sup>ST</sup>	L	E	A	N	A	A
	2 <sup>ND</sup>	N	N	E	A	N	N

The intensivist is prompted next to determine, sequentially, table input values for CAD, RAD, and EF.

In these decision tables, the letter references have the following meanings: L=labetalol, E=esmolol, A=enalapril, N=nicardipine, H=hydralazine, S=nitroprusside. The reference to 1<sup>st</sup>

1 and 2<sup>nd</sup> means that treatment should begin with the 1<sup>st</sup> drug and add or substitute the 2<sup>nd</sup> drug as  
2 needed.

3 Using the above decision tables, the intensivist is prompted to determine whether the  
4 patient has known coronary artery disease (CAD) or 3 or more risk factors for CAD **1824**. If this  
5 criteria is met **1824**, the intensivist is prompted to enter a “Y” or “YES” for CAD into the table  
6 selected above in **1820** and **1822**. If this criteria is not met, the intensivist is prompted to enter a  
7 “N” or “NO” for CAD into the table selected above in **1820** and **1822**.

8 Next, the intensivist is prompted to determine whether the patient has known reactive  
9 airway disease (RAD)**1826**. If this criteria is met **1826**, the intensivist is prompted to enter a “Y”  
10 or “YES” for RAD into the table selected above in **1820** and **1822**. If this criteria is not met, the  
11 intensivist is prompted to enter a “N” or “NO” for RAD into the table selected above in **1820** and  
12 **1822**.

13 Next, the intensivist is prompted to determine whether the patient has known EF less than  
14 30 percent or a history of systolic heart failure **1828**. If this criteria is met **1828**, the intensivist is  
15 prompted to enter a “Y” or “YES” for EF into the table selected above in **1820** and **1822**. If this  
16 criteria is not met **1828**, the intensivist is prompted to enter a “N” or “NO” for EF into the table  
17 selected above in **1820** and **1822**.

18 Based on the table selected in **1820** and **1822** above, and the table inputs determined from  
19 **1824**, **1826**, and **1828**, the intensivist is prompted with the proper medication to administer for  
20 the 1<sup>st</sup> and 2<sup>nd</sup> treatment.

21 If the patient is not in pain **1812**, the intensivist is prompted to employ the procedures  
22 described above in **1816**.



perform a VQ scan and perform further assessment of the patient **2904**.

In order to further assess the patient, the intensivist is prompted to respond to a series of questions **2906, 2908, 2910, 2912**. The intensivist is prompted to determine whether any of the following patient conditions are present: Dyspnea, Worsening chronic dyspnea, Pleuritic chest pain, Chest pain that is non- retro sternal & non- pleuritic, O<sub>2</sub> saturation < 92% on room air that corrects with 40% O<sub>2</sub> supplementation, Hemoptysis, or Pleural rub **2906**. The intensivist is also prompted to determine whether any risk factors are in the patient's history, such as: Surgery within 12 weeks, Immobilization (complete bed rest) for > 3 days within 4 weeks, Previous DVT or objectively diagnosed PE, Lower extremity fracture & immobilization within 12 weeks, Strong family history of DVT or PE (  $\geq 2$  family members with objective proven events or 1<sup>st</sup> degree relative with hereditary thrombophilia), Cancer (treatment within the last 6 months or palliative stages), Postpartum, or Lower extremity paralysis **2908**. Further, the intensivist must determine whether the patient has any of the following symptoms: Heart rate > 90 beats/min, Temp  $\geq 38.0$ , CXR free of abnormalities (edema, pneumonia, pneumothorax), or Leg symptoms c/w DVT, syncope, blood pressure less than 90 mm Hg with heart rate greater than 100 beats/min, receiving mechanical ventilation and/or oxygen supplementation greater than 40%, and new onset or right heart failure (-JVP, new S1, Q3, T3, or RBBB) **2910**. The intensivist is also queried by the system to consider alternative diagnosis that may be more likely than pulmonary embolism. To do so, the intensivist is prompted to consider conditions that simulate major pulmonary embolism, such as myocardial infarction, acute infection with COPD, septic Shock, dissecting aortic aneurysm, or occult hemorrhage. The intensivist is additionally prompted to consider conditions that simulate minor pulmonary embolism, such as acute

1 bronchitis, pericarditis, viral pleurisy, pneumonia, and esophageal spasm **2912**.

2 Referring to **Figure 31A**, the pulmonary embolism algorithm description continues. The  
3 intensivist enters the answers to the assessment queries posed **2906, 2908, 2910, 2912** into the  
4 system. If two or more responses to the patient condition query **2906** were answered yes and one  
5 or more questions were answered yes from: Heart rate  $> 90$  beats/min, Temp  $\geq 38.0$ , CXR free of  
6 abnormalities, or Leg symptoms c/w DVT of the symptoms query **2910**, the intensivist is  
7 informed that a typical pulmonary embolism is present **2914**. Next, the system compares this  
8 response to the answer to the alternative diagnosis query **2912**. If an alternative diagnosis is at  
9 least as likely as pulmonary embolism **2916**, the intensivist is also given a low probability **2918**  
10 to moderate probability **2920** risk factor. If an alternative diagnosis is less likely than pulmonary  
11 embolism **2922**, the intensivist is given a moderate **2924** to high **2926** probability risk factor.

12 If less than two yes answers resulted from the patient conditions **2906**, the intensivist is  
13 advised by the system that an atypical pulmonary embolism may be present **2928**. Next, the  
14 system compares this response to the answer to the alternative diagnosis query **2912**. If an  
15 alternative diagnosis is at least as likely as pulmonary embolism **2930**, the intensivist is told there  
16 is no risk and low probability **2932** or some risk with a low probability **2934** risk factor. If an  
17 alternative diagnosis is less likely than pulmonary embolism **2934**, the intensivist is given a no  
18 risk and low probability **2938** to risk but moderate probability **2940**.

19 If at least one answer to the symptoms of syncope, blood pressure less than 90 mm Hg  
20 with heart rate greater than 100 beats/min, receiving mechanical ventilation and/or oxygen  
21 supplementation greater than 40%, and new onset or right heart failure **2910** is yes, the  
22 intensivist is prompted with a message that severe pulmonary embolism is occurring **2942**. Next,



the system compares this response to the answer to the alternative diagnosis query **2912**. If an alternative diagnosis is at least as likely as pulmonary embolism **2944**, the intensivist is told there is a moderate probability of pulmonary embolism **2946**. If an alternative diagnosis is less likely than pulmonary embolism **2948**, the intensivist is notified that a high probability of pulmonary embolism is present **2950**.

Once the risk factors and probabilities are determined the system compares this information to the VQ scan results. This comparison is performed according to the following Table 4 below.

**Table 4: Probability table**

<u>V/Q Scan</u>	<u>Clinical Probability</u>		
	High	Moderate	Low
High	A	A	B
Intermediate	B	C	C
Low	B	C	E
Normal	E	E	E

Where the VQ scan column and the risk column intersect, a letter code is assigned to various treatment instructions. The treatment instructions are as follows.

**A = Pulmonary embolus diagnosed. Begin treatment**

E = Pulmonary embolus excluded

B = Proceed with the following work-up:

- 1) Perform spiral CT(If patient has renal insufficiency [creatinine > 2.0], consider going directly to pulmonary angiogram to reduce the potential dye load). If positive begin treatment,

- 2) If negative, assess for DVT using compression ultrasound or venography. If positive begin treatment,
- 3) If negative, perform pulmonary angiogram. If positive begin treatment, if negative diagnosis excluded.

C = Proceed with the following work-up:

- 1) Perform spiral CT. If positive begin treatment,
- 2) If negative, assess for DVT using compression ultrasound or venography. If positive begin treatment,
- 3) If negative perform D-dimer assay(elisa only). If negative diagnosis excluded, If positive, perform serial ultrasound of the lower extremities.

Once the correlation is made, the instructions associated with the letter code are displayed by the system to prompt the intensivist with diagnosis and treatment instructions.

Referring to **Figure 32**, the seizure decision support algorithm of the present invention is illustrated. If an intensivist encounters seizure in a patient, he may not be certain of all of the aspects and the timelines that are critical to treating this particular condition. Therefore, the intensivist is lead through a decision support algorithm, which divides the treatment sequence into three segments: 0-30 minutes; 30-60 minutes; and beyond 60 minutes.

At the onset of a seizure, in the 0-30 minute segment of the algorithm, the intensivist is prompted to give the patient lorazepam (0.1 mg/kg of bodyweight) in 2 mg boluses up to 8 mg **2000**. Subsequently, the intensivist is prompted to give the patient phenytoin (18-20 mg/kg of bodyweight) at 50mg/min of fosphenytoin (18-20 mg/kg of bodyweight) at 150 mg/min followed by 5 mg/kg of bodyweight/day through separate IV line **2002**.

During the 30-60 minute segment of the algorithm, the intensivist is prompted to: reload additional phenytoin or fosphenytoin (10 mg/kg of bodyweight) maintaining previous infusion; and give additional lorazepam (0.05 mg/kg of bodyweight) **2004**. Subsequently, the intensivist is prompted to begin continuous EEG monitoring **2006**.

1 The intensivist is then prompted to determine whether the patient is hemodynamically  
2 stable **2008**. If hemodynamically stable, the intensivist is prompted to administer propofol 1-2  
3 mg/kg of bodyweight bolus followed by 2-10 mg/kg/hr **2010**.

4 At the 60 minute segment of the algorithm, the intensivist is prompted that if seizure  
5 activity stops, he should taper either midazolam or propofol over the next 12-24 hours while  
6 maintaining phenytoin but if seizures persist, he is prompted to move to the pentobarbital coma  
7 block **2012**.

8 Under pentobarbital coma, the intensivist is prompted to administer 10-15 mg/kg/hr and  
9 to maintain until seizure control is achieved on EEG 2014. The intensivist is prompted further  
10 that the patient usually requires PA catheter and pressors to maintain hemodynamic control **2014**.

11 Alternatively, if the patient is determined to be hemodynamically unstable **2016**, the  
12 intensivist is prompted to utilize fluids and pressors as needed (phynylephrine or dopamine)  
13 midazolam 0.2 mg/kg bolus followed by 0.1-2.0 mg/kg/hr 2018.

14 At the 60 minute segment of the algorithm, the intensivist is prompted that if seizure  
15 activity stops, he should taper either midazolam or propofol over the next 12-24 hours while  
16 maintain phenytoin but if seizures persist, he is prompted to move to the pentobarbital coma  
17 block **2012**.

18 Under pentobarbital coma, the intensivist is prompted to administer 10-15 mg/kg/hr and  
19 to maintain until seizure control is achieved on EEG 2014. The intensivist is prompted further  
20 that the patient usually requires PA catheter and pressors to maintain hemodynamic control **2014**.

Referring to **Figure 33**, the supra ventricular tachycardia (SVT) decision support algorithm of the present invention is illustrated. If an intensivist determines that SVT is present, the intensivist may not be certain of all aspects that would be involved in treating this particular condition. Therefore, the intensivist is lead through a decision support algorithm which prompts the intensivist to determine the appropriate care to be given.

Initially, the intensivist is prompted to determine whether SVT is stable or unstable **2100**. If SVT is stable **2102**, the intensivist is prompted to determine whether the patient has a regular or irregular rhythm **2102**. If the patient has a regular rhythm **2104**, the intensivist is prompted to determine whether there is a wide complex or a narrow complex **2104**. If the intensivist determines that there is a wide complex **2106**, the intensivist is prompted to administer adenosine 6 mg/12 mg (if needed) **2108**. Following the administering of adenosine **2108**, the intensivist is prompted to consider that if the patient converts to sinus rhythm (SR) to – consider re-entrant junctional or WPW re-entrant. If the wide complex recurs, treat the patient with esmolol or Ca+2 blockers.

Alternatively; if no effect, the intensivist is prompted to consider V-tach **2112**. Next, the intensivist is prompted to: 1) load procainamide 150 mg over 10 min, then 1 mg/min infusion; and 2) synchronized cardiovert **2114**.

Alternatively, if the wide complex slows, the intensivist is prompted to consider SVT w/ aberrancy and continue to slow with esmolol or Ca+2 blockers **2116**.

The intensivist is prompted next to administer esmolol/calcium blockers and link to ventricular rate control **2118**. The intensivist is prompted next to determine whether there has been a conversion to SR **2120**. If there is no conversion to SR in 24 hours, the intensivist is

1 prompted to add antiarrhythmic agent and consider anticoagulation **2122**. The intensivist is  
2 prompted next to determine whether there has been conversion to SR. If conversion to SR, the  
3 intensivist is prompted to continue maintenance antiarrhythmic agent during hospitalization  
4 **2124**. If no conversion to SR, the intensivist is prompted to cardiovert while on antiarrhythmic  
5 & following heparinization **2126**.

6 If the patient has a regular rhythm **2104**, the intensivist is prompted to determine whether  
7 there is a wide complex or a narrow complex **2104**. If the intensivist determines that there is a  
8 narrow complex **2128**, the intensivist is prompted to to administer adenosine 6mg/12mg (if  
9 needed) **2130**. If administering the adenosine **2130** slows the ventricular rate only and the atrial  
10 rate persists, the intensivist is prompted to consider atrial flutter and continue to slow with  
11 esmolol or Ca+2 blockers **2132**. The intensivist is prompted next to employ the procedures  
12 described above in **2118**.

13 If administering the adenosine **2130** converts the patient to SR, the intensivist is  
14 prompted to consider re-entrant sinus or junctional and if recurs, treat with esmolol or Ca+2  
15 blockers **2134**.

16 If administering the adenosine **2130** slows both atrial and ventricular rates the intensivist  
17 is prompted that there is a probable sinus tachycardia **2136**. The intensivist is prompted next to  
18 continue to slow with esmolol **2138**. The intensivist is prompted next to employ the procedures  
19 described above in **2118**.

20 If SVT is stable **2102**, the intensivist is also prompted to determine whether the patient  
21 has a regular or irregular rhythm **2102**. If the patient has an irregular rhythm **2140**, the  
22 intensivist is prompted that if no p waves, there is probable Atrial fibrillation **2142**. The

1 intensivist is prompted next to slow ventricular response with esmolol or Ca+2 blockers **2144**.

2 The intensivist is prompted next to employ the procedures described above in **2118**.

3 If the patient has an irregular rhythm **2140**, the intensivist is prompted to determine  
4 whether there are more than 3 p wave types MAT – and to treat underlying lung dz. and avoid  
5 theophylline compounds **2146**. The intensivist is prompted next to slow rate with Ca+2 blockers  
6 only **2148**. The intensivist is prompted next to employ the procedures described above in **2118**.

7 Referring now to **Fig. 33A**, the description of the SVT decision algorithm continues. If  
8 SVT is unstable **2101**, the intensivist is prompted to determine whether the patient has SBP less  
9 than 80, ischemia, mental status changes **2150**. The intensivist is prompted next to perform  
10 synchronous cardioversion (100 J, 200 J, 300 J) **2152**. The intensivist is prompted next that if  
11 sinus rhythm: 1) correct reversible etiologies; 2) consider starting IV antiarrhythmic for  
12 maintenance of sinus rhythm **2154**. Alternatively, following **2152**, the intensivist is prompted  
13 next that if continued SVT: 1) correct reversible etiologies; 2) load IV antiarrhythmic (see dosing  
14 guidelines) and repeat DC cardioversion **2156**.

15 For example, and without limitations, wide complex QRS Tachycardia is also addressed  
16 in the decision support algorithm of the present invention. Referring to **Figure 34**, the wide  
17 complex QRS tachycardia decision support algorithm is illustrated. If an intensivist determines  
18 that there may be a possibility of wide complex QRS tachycardia, the intensivist may not be  
19 certain of all aspects that would be involved in this particular condition. Therefore, the  
20 intensivist is lead through a decision support algorithm which prompts the intensivist to  
21 determine the appropriate care to be given.

22 Initially, the intensivist is prompted to determine whether the patient is hemodynamically

1 stable (no angina, heart failure, or hypotension (systolic less than 80 mm)) **2200**. If this criteria  
2 is not met, the intensivist is prompted to go to the cardio-pulmonary guidelines algorithm which  
3 is generally known to those skilled in the art.

4 Alternatively, if this criteria is met, the intensivist is prompted to determine whether the  
5 patient is within 7 days of a myocardial infarction or at risk for myocardial ischemia **2202**. If the  
6 patient is not within 7 days of a myocardial infarction or at risk for myocardial ischemia **2202**,  
7 the intensivist is prompted to determine whether the wide complex QRS rhythm is sustained  
8 (greater than 30 seconds) **2234**. If this criteria is not met, the intensivist is prompted to  
9 determine whether the QRS is monomorphic **2236**. If the QRS is monomorphic **2236**, the  
10 intensivist is prompted to determine whether the patient has structural heart disease **2242**. If the  
11 patient has structural heart disease **2242**, the intensivist is prompted to: 1) monitor closely; 2)  
12 look for reversible etiologies; and 3) consider antiarrhythmic therapy **2244**. If the patient does  
13 not have structural heart disease **2242**, the intensivist is prompted to: 1) monitor closely; 2) look  
14 for reversible etiologies; and 3) if recurs and symptomatic may require further testing (prolonged  
15 holter or EP study) **2246**.

16 If the QRS is not monomorphic **2236**, the intensivist is prompted to determine whether  
17 the QT is prolonged **2238**. If this criteria is met, the intensivist is prompted to: 1) check K; 2)  
18 give Mg; and 3) consider overdrive pacing **2240**. If the intensivist determines that the QT is not  
19 prolonged, **2238**, the intensivist is prompted to employ the procedures described above in **2242**.

20 If the wide complex QRS rhythm is sustained **2234**, the intensivist is prompted to  
21 determine whether the rhythm is polymorphic or irregular **2208**. If the rhythm is polymorphic or  
22 irregular, the intensivist is prompted to consider atrial fibrillation with accessory pathway

1 conduction and load with procainamide and get a cardiology consultation **2210**. If the rhythm is  
2 not polymorphic or irregular, the intensivist is prompted with the question of whether he wishes  
3 to: 1) perform ECG diagnosis; or 2) administer adenosine diagnostically **2220**. If the intensivist  
4 makes the determination to perform an ECG diagnosis **2220**, he is prompted to go to the ECG  
5 diagnosis algorithm **2300**.

6 If the intensivist makes the determination to administer adenosine diagnostically **2220**, he  
7 is prompted to go to the administer adenosine branch of the algorithm **2222**. If there is no effect,  
8 the intensivist is prompted that there is probable VT and to determine whether the VT is  
9 monomorphic **2224**. If the VT is monomorphic **2224**, the intensivist is prompted to load with  
10 procainamide and perform synchronous cardioversion **2226**.

11 Alternatively, if the VT is not monomorphic **2224**, the intensivist is prompted to load  
12 with lidocaine and perform immediate cardioversion **2228**.

13 If the ventricular response is slowed after administering adenosine **2222**, the intensivist is  
14 prompted to consider SVT with aberrancy and treat with esmolol or Ca blockers **2230**.

15 If the ventricular response converts to sinus rhythm after administering adenosine **2222**,  
16 the intensivist is prompted: to consider re-entrant mechanism with BBB or WPW; and, 1) if  
17 WPW consult cardiology for possible ablation **2232**.

18 If the patient is within 7 days of a myocardial infarction or at risk for myocardial  
19 ischemia **2202**, the intensivist is prompted to determine whether the wide complex is sustained  
20 (30 seconds) **2204**. If the wide complex is not sustained **2204**, the intensivist is prompted to  
21 determine whether the patient: 1) symptomatic; 2) tachycardia runs are frequent; or 3) the  
22 tachycardia rates are rapid (greater than 180) **2212**. If this criteria is not met, the intensivist is



1 prompted to observe **2216**. Alternatively, if this criteria is met **2212**, the intensivist is prompted  
2 to: 1) administer lidocaine 100-200 mg & 1-4 mg/min infusion; and 2) amiodarone **2214**.

3 If the wide complex is sustained **2204**, the intensivist is prompted to determine whether  
4 the rate is greater than 140/min **2206**. If this criteria is not met **2206**, the intensivist is prompted:  
5 to consider accelerated idioventricular, and that in some patients this can lead to hemodynamic  
6 compromise; and that 1) he can perform overdrive pacing if needed **2218**.

7 Alternatively, if this criteria is met, the intensivist is prompted to follow the procedures in  
8 **2208**.

9 If the intensivist makes the determination to perform ECG Diagnosis **2220**, he is  
10 prompted to go to the ECG Diagnosis branch of the algorithm **2220**. Referring now to Figure  
11 34A, in the ECG Diagnosis branch, the intensivist is prompted to determine whether the patient  
12 has known pre-excitation syndrome **2300**. If this criteria is met, the intensivist is prompted to  
13 determine whether the QRS complexes are predominantly negative in leads V4-V6 **2302**. If the  
14 QRS complexes are predominantly negative in leads V4-V6, the intensivist is prompted that  
15 there is probable VT **2304**.

16 If the QRS complexes are not predominantly negative in leads V4-V6 **2302**, the  
17 intensivist is prompted to determine whether there is a QR complex in one or more of leads V2-  
18 V6 **2306**. If this criteria is met, the intensivist is prompted that there is probable VT **2308**.

19 Alternatively, if this criteria is not met **2306**, the intensivist is prompted to determine  
20 whether there are more QRS complexes than P waves **2310**. If there are more QRS complexes  
21 than P waves **2310**, the intensivist is prompted that there is probable VT **2312**. If there are not  
22 more QRS complexes than P waves **2310**, the intensivist is prompted: to consider pre-excited

SVT; and that he may wish to perform EP study **2314**.

If the intensivist determines that the patient does not have known pre-excitation syndrome **2300**, the intensivist is prompted to determine whether there is an RS complex present in any precordial lead **2316**. If this criteria is not met **2316**, the intensivist is prompted that there is probable VT **2318**.

Alternatively, if this criteria is met **2316**, the intensivist is prompted to determine whether the R to S interval is greater than 100 MS in any one precordial lead **2320**. If this criteria is met, the intensivist is prompted that there is probable VT **2322**.

If the R to S interval is not greater than 100 MS in any one precordial lead **2320**, the intensivist is prompted to determine whether there is evidence of atrioventricular dissociation **2324**. If this criteria is met, the intensivist is prompted that there is probable VT **2326**.

Alternatively, if there is no evidence of atrioventricular dissociation **2324**, the intensivist is prompted to determine whether V-1 is negative and V-6 positive and QRS greater than 0.14 mSEC **2328**. If this criteria is met, the intensivist is prompted that there is probable VT **2330**.

If this criteria is not met **2328**, the intensivist is prompted that the situation may represent SVT with aberrancy or underlying bundle branch block **2332**.

Referring to **Figure 41**, the assessment of sedation algorithm of the present invention is illustrated. If an intensivist encounters a need for sedation, he may not be certain of all of the aspects and the timelines that are critical to this particular process. Therefore, the intensivist is lead through a decision support algorithm, which prompts the intensivist to address a number of factors in the process **3100**.

1 The intensivist is prompted initially to go to the Scoring section of the algorithm **3100**.  
2 The intensivist is prompted to proceed through a number of scorings **3102** and to first score the  
3 patient's alertness with points being allocated in the following manner: asleep/unresponsive=0;  
4 responsive to voice=1; and hyperresponsive=2 **3104**.

5 The intensivist is prompted next to score the patient's movement with points being  
6 allocated in the following manner: no spontaneous movement=0; spontaneous movement=1; and  
7 pulls at lines, tubes, dressings=2 **3106**.

8 The intensivist is prompted next to score the patient's respiration based on whether the  
9 patient is mechanically ventilated or spontaneously breathing with points being allocated as  
10 subsequently discussed. If the patient is mechanically ventilated, the intensivist is prompted to  
11 allocate points in the following manner: no spontaneous ventilation=0; spontaneous ventilations  
12 and synchronous with ventilator=1; or spontaneous ventilations with cough or dysynchrony>10  
13 percent of breaths=2 **3108**. Alternatively, if the patient is spontaneously breathing, the  
14 intensivist is prompted to allocate points in the following manner: respiration rate (RR) <10=0;  
15 RR=10-30=1; or RR>30=2 **3108**.

16 The intensivist is prompted next to score the patient's heart rate with points being  
17 allocated in the following manner: >20 percent below mean for last 4 hr=0; within 20 percent  
18 mean for last 4 hr=1; or >20 percent above mean for last 4 hr=2 **3110**.

19 The intensivist is prompted next to score the patient's blood pressure with points being  
20 allocated in the following manner: MAP >20 percent for last 4 hr=0; MAP within 20 percent  
21 mean for last 4 hr=1; or MAP >20 percent above mean for last 4 hr=2 **3112**.

1           The intensivist is prompted next to determine the sedation score by the following  
2   formula: SEDATION SCORE=alertness + movement + respirations + heart rate + blood  
3   pressure **3114**. In one embodiment, respiratory rate, heart rate, and BP can be computer linked to  
4   monitor data thereby simplifying the sedation scoring assessment. The nursing observations are  
5   deemed intuitive and the nursing burden in sedation scoring can be minimal by using this point  
6   scoring.

7           Referring now to **Figure 41A**, the sedation assessment algorithm description continues.  
8   The intensivist is prompted then to continue the sedation assessment by moving to the Pain  
9   Assessment section of the algorithm **3116**.

10           In the Pain Assessment section, the intensivist is prompted to determine whether the  
11   patient is conscious, communicative, and acknowledging pain **3118**. If this criteria is not met,  
12   the intensivist is prompted to determine: whether the sedation score is greater than 2 and the  
13   patient: is known to be in pain before becoming uncommunicative; or S/p recent surgery; or  
14   having tissue ischemia or infarct; or has wounds; or has large tumor possibly impinging on  
15   nerves. If the answer to either of these two questions is YES, the intensivist is prompted to treat  
16   for pain **3118**. The intensivist is prompted then to continue the assessment by moving to the  
17   Delirium Assessment section of the algorithm **3118**.

18           In the Delirium Assessment section, the intensivist is prompted to determine whether the  
19   sedation score is greater than 2 AND the patient has: day/night reversal with increased agitation  
20   at night OR eyes open and “awake” but disoriented; or eyes open and “awake” but pulling at  
21   lines, tubes, or dressings OR difficult to sedate prior to ventilator weaning OR paradoxical

1 response to benzodiazepines. If this criteria is met, the intensivist is prompted to consider  
2 butyrophenone **3120**.

3 Referring to **Figure 42**, the Bolus sliding scale algorithm is illustrated. If an intensivist  
4 encounters a need for sedation, the algorithm for which may contain a reference to the bolus  
5 sliding scale for midazolam, he may not be certain of all of the aspects which are critical to this  
6 scale. Therefore, the intensivist is lead through a decision support algorithm, which prompts the  
7 intensivist through the use of the scale **3200**.

8 If lorazepam is less than 0-2 mg IV q 6hr, then the intensivist is prompted to give  
9 midazolam 1-2 mg q 5 min until adequately sedated **3202**.

10 Alternatively, if lorazepam equals 2-4 mg IV q 4 hr, then the intensivist is prompted to  
11 give midazolam 2 mg q 5 min until adequately sedated **3202**.

12 Alternatively, if lorazepam is greater than 10 mg IV q 4 hr, then the intensivist is  
13 prompted to give midazolam 5 mg q 5 min until adequately AND consider fentanyl and/or  
14 droperidol or Haldol for synergy despite delirium and pain assessment **3202**.

15 Yet another decision support routine is the sedation algorithm. Referring to **Figure 43**,  
16 the sedation process decision support algorithm is illustrated. If an intensivist determines that a  
17 patient will require sedation, the intensivist may not be certain of all aspects that would be  
18 involved in this particular process. Therefore, the intensivist is lead through a decision support  
19 algorithm, which prompts the intensivist to conduct a sedation assessment based on: 1) scoring;  
20 2) pain; and 3) delirium (see Assessment of Sedation algorithm) **3300**.

21 Following completion of the sedation assessment process **3300**, the intensivist is  
22 prompted to determine whether the patient is in pain **3302**. If this criteria is met, the intensivist

1 is prompted to administer bolus morphine, fentanyl, other narcotic, start patient controlled  
2 analgesic (PCA) or epidural analgesia as indicated **3324**. If the patient is not in pain **3302** or  
3 after administering bolus morphine, fentanyl, other narcotic, start patient controlled analgesic  
4 (PCA) or epidural analgesia as indicated **3324**, the intensivist is prompted to determine whether  
5 the patient is delirious **3304**.

6 If the intensivist determines that the patient is delirious **3304**, he is prompted to  
7 administer droperidol 2.5-5 mg q30min prn and that he may consider IV Haldol not to exceed  
8 30mg/24hr **3326**. If the patient is not delirious or after following the procedures in **3326**, the  
9 intensivist is prompted to determine whether the patient will need sedation for more than the next  
10 24 hours **3306**. If the patient will not need sedation for more than the next 24 hours **3306**, the  
11 process continues as described in **Figure 44**.

12 Alternatively, if the patient will need sedation for more than the next 24 hours **3306**, the  
13 intensivist is prompted to determine whether the sedation score is 8-10 **3308**. If this criteria is  
14 met, the intensivist is prompted to employ the Bolus sliding scale midazolam and increase  
15 lorazepam by 20 percent **3328** (see Bolus sliding scale midazolam algorithm – **Figure 42**).  
16 Subsequently, the intensivist is prompted to reassess sedation in 4 hr **3330**.

17 If the sedation score is not 8-10, the intensivist is prompted to determine whether the  
18 sedation score is greater than or equal to the last Sed Scr after sedative bolus or increase **3310**. If  
19 this criteria is met, the intensivist is prompted to employ the procedures described above in **3328**  
20 and **3330**.

21 If the sedation score is not greater than or equal to the last Sed Scr after sedative bolus or  
22 increase **3310**, the intensivist is prompted to determine whether four (4) or more midaz boluses

1 have been given since last q4hr assessment **3312**. If this criteria is met, the intensivist is  
2 prompted to employ the procedures described above in **3328** and **3330**.

3 Alternatively, if less than four (4) midaz boluses have been given since last q4hr  
4 assessment **3312**, the intensivist is prompted to determine whether the patient is adequately  
5 sedated **3314**. If this criteria is not met, the intensivist is prompted to employ the procedure  
6 described in **3328** and **3330**.

7 If the intensivist determines that the patient is adequately sedated **3314**, the intensivist is  
8 prompted to determine whether the sedation score is 0-2 **3316**. If this criteria is met, the  
9 intensivist is prompted to decrease lorazepam by 20 percent **3332** and reassess sedation in 4 hr  
10 **3334**.

11 Alternatively, if the sedation score is not 0-2 **3316**, the intensivist is prompted to  
12 determine whether the sedation score is less than or equal to the last Sed Scr after sedative  
13 decrease **3318**. If this criteria is met, the intensivist is prompted to employ the procedure  
14 described in **3332** and **3334**.

15 If the sedation score is not less than or equal to the last Sec Scr after sedative increase  
16 **3318**, the intensivist is prompted to determine whether the patient is clinically oversedated **3320**.

17 If the patient is clinically oversedated **3320**, the intensivist is prompted to employ the procedure  
18 described in **3332** and **3334**. If the patient is not clinically oversedated **3320**, the intensivist is  
19 prompted to reassess sedation in 4 hr **3322**.

20 Referring to **Figure 44**, the short term sedation process decision support algorithm of the  
21 present invention is illustrated. If an intensivist determines that a patient will not require  
22 sedation past the next 24 hour period, the intensivist may not be certain of all aspects that would

1 be involved in this particular process. Therefore, the intensivist is lead through a decision  
2 support algorithm, which prompts the intensivist to conduct a sedation assessment based on: 1)  
3 scoring; 2) pain; and 3) delirium (see Assessment of Sedation algorithm) **3100**.

4 Following completion of the sedation assessment process **3100**, the intensivist is  
5 prompted to decrease lorazepam by 20 percent from baseline per day **3102**. The intensivist is  
6 prompted next to determine whether the patient is in pain **3104**. If this criteria is met, the  
7 intensivist is prompted to administer bolus morphine or fentanyl **3122**. If the patient is not in  
8 pain or after administering bolus morphine or fentanyl **3122**, the intensivist is prompted to  
9 determine whether the patient is delirious **3106**.

10 If the intensivist determines that the patient is delirious, he is prompted to administer  
11 droperidol 2.5-5 mg q30min prn **3124**. If the patient is not delirious or after administering  
12 droperidol **3124**, the intensivist is prompted to determine whether the sedation score is 8-10  
13 **3108**.

14 If this criteria is met, the intensivist is prompted to employ the Bolus sliding scale  
15 midazolam (see Bolus sliding scale midazolam algorithm) and begin midazolam infusion or  
16 begin propofol 1-2 mg/kg bolus and 5-50 mcg/kg/min infusion **3126**. Subsequently, the  
17 intensivist is prompted to reassess sedation in 1 hr **3128**.

18 If the sedation score is not 8-10, the intensivist is prompted to determine whether the  
19 sedation score is greater than or equal to the last Sed Scr after sedative bolus or increase **3110**. If  
20 this criteria is met, the intensivist is prompted to employ the procedures described above in **3126**  
21 and **3128**.

22 If the intensivist determines that the sedation score is not greater than the last sedation



1 score after sedative bolus or increase **3110**, the intensivist is prompted to determine whether the  
2 patient is adequately sedated **3112**. If this criteria is not met, the intensivist is prompted to  
3 employ the procedures described above in **3126** and **3128**.

4 If the intensivist determines that the patient is adequately sedated **3112**, he is prompted to  
5 determine whether the sedation score is 0-2 **3114**. If this criteria is met, the intensivist is  
6 prompted to determine if the patient has been sedated for more than 72 hours **3130**. If the  
7 patient has not been sedated for more than 72 hours **3130**, the intensivist is prompted to hold  
8 midazolam or propofol and hold or decrease lorazepam by 50 percent **3132**. The intensivist is  
9 prompted subsequently to reassess sedation in 1 hour **3134**.

10 Alternatively, if the intensivist determines that the patient has been sedated for more than  
11 72 hours **3130**, the intensivist is prompted to hold midazolam or propofol and decrease  
12 lorazepam by 20 percent per day **3136**. The intensivist is prompted subsequently to reassess  
13 sedation in 1 hour **3134**.

14 Alternatively, if the intensivist determines that the sedation score is not 0-2 **3114**,  
15 the intensivist is prompted to determine whether the sedation score is less than or equal to the  
16 last sedation screening after sedative decrease **3116**. If this criteria is met, the intensivist is  
17 prompted to determine whether the patient has been sedated for more than 72 hours and to  
18 follow the procedures described above in **3130**.

19 If the intensivist determines that the sedation score is not less than or equal to the  
20 last Sed Scr after sedative decrease **3116**, the intensivist is prompted to determine whether the  
21 patient is clinically oversedated **3118**. If this criteria is met, the intensivist is prompted to  
22 determine whether the patient has been sedated for more than 72 hours and to follow the

1 procedures described above in **3130**. If this criteria is not met, the intensivist is prompted to  
2 reassess sedation in 1 hr **3120**.

3 Referring to **Figure 45**, the respiratory isolation decision support algorithm is illustrated.  
4 If an intensivist determines that there may be a need for respiratory isolation, the intensivist may  
5 not be certain of all aspects that would be involved in this process. Therefore, the intensivist is  
6 lead through a decision support algorithm which prompts the intensivist to determine the need  
7 for respiratory isolation based upon: a) clinical assessment; and/or b) smear/culture findings  
8 **3500**.

9 Pursuing the clinical assessment branch of the decision support algorithm, the intensivist  
10 is prompted to determine whether the patient has known mTB (mycobacterium tuberculosis)  
11 **3502**. If this criteria is met, the intensivist is prompted to determine whether the patient has been  
12 compliant with their medications for over 2 weeks and is clinically responding **3512**. If the  
13 patient has not been compliant with their medications for over 2 weeks and is not clinically  
14 responding **3512**, the intensivist is prompted that isolation is required **3514**. If the patient has  
15 been compliant with their medications and is clinically responding **3512**, the intensivist is  
16 prompted that no isolation is required **3516**.

17 Alternatively, if the patient does not have known mTB **3502**, the intensivist is prompted  
18 to determine whether the patient has known mycobacterial disease other than TB **3504**. If this  
19 criteria is met, the intensivist is prompted to determine whether the patient has new CXR (chest x  
20 ray) findings and symptoms (cough 2 weeks, fever, weight loss) **3518**. If the patient does not  
21 have new CXR findings and symptoms **3518**, the intensivist is prompted that no isolation is

required **3520**. If the patient does have new CXR findings and symptoms **3518**, the intensivist is prompted that isolation is required **3522**.

If the patient does not have known mycobacterial disease other than TB **3504**, the intensivist is prompted to determine whether there is a new cavitory lesion on CXR **3506**. If this criteria is met, the intensivist is prompted that isolation is required **3524**.

Alternatively, if there is no new cavitory lesion on CXR **3506**, the intensivist is prompted to determine whether there are pulmonary infiltrates or whether the patient is HIV (human immunodeficiency virus) positive **3508**. If this criteria is not met, the intensivist is prompted that no isolation is required **3510**. If this criteria is met, the intensivist is prompted to determine whether the patient has new CXR findings and symptoms (cough 2 weeks, fever, weight loss) **and** at high risk: 1) known mTB exposure; 2) homeless; 3) prisoner; 4) travel to area with multi-drug resistant TB **3526**. If this criteria is met, the intensivist is prompted that isolation is required **3528**. Alternatively, if this criteria is not met, the intensivist is prompted that no isolation is required **3530**.

Pursuing the smear/culture branch of the decision support algorithm **3500**, the intensivist is prompted to determine whether the AFB (acid-fast bacilli) smear is positive **3532**. If the AFB smear is not positive, the intensivist is prompted that: no isolation is required; await culture results; if culture negative, no isolation required; if culture positive and patient has mycobacterial disease other than TB (MOTT no isolation is required; if the culture is positive and the patient does not have MOTT consult ID **3534**.

Alternatively, if the AFB smear is positive, the intensivist is prompted to determine whether the patient has known mycobacterial disease other than TB **3536**. If this criteria is not

met, the intensivist is prompted that isolation is required **3538**. If this criteria is met, the intensivist is prompted: to isolate until results of NAP test are in; if mTB positive isolate the patient; if no mTB, no isolation is required **3540**.

Referring to **Figure 47**, the empiric meningitis treatment decision support algorithm of the present invention is illustrated. If the intensivist is treating a patient for meningitis, the intensivist is prompted to answer a series of queries by the system to properly address medication and dosage. First, the intensivist is prompted to determine whether the patient has suffered a head trauma or undergone neurosurgery **3700**. The answer to this question is input 1 to **table x** below. The intensivist is next prompted to determine whether the patient is allergic to penicillin or is from an area where penicillin resistant staphylococcus pneumoniae is prevalent **3702**. The answer to this question becomes input 2 to **table x** below. The intensivist must also determine whether the patient is immunocompromised **3704**, and the answer becomes input 3 to **table x** below. The intensivist determines if the patient is over fifty years of age **3706**, with the answer being input 4 in **table x** below. Lastly, the intensivist is prompted to determine whether the patient has altered mental status **3708**, and the answer becomes input 5 in **table x** below. The inputs to each of these prompts **3702, 3704, 3706, 3708** is compared to a dosage database according to the **Table 5** below.

**Table 5:** Meningitis Input-Output Table

Input	Combinations	Output
1	1 = yes 2 = no	A) vancomycin 1.5 – 2 gm IV q 12h + ceftazidone 2gm IV q 8 hr or cefapime 2gm IV q 8 hr
2	1 = yes 2 = no	B) vancomycin 1.5 – 2 gm IV q 12h

		+ aztreonam 0.5 – 2 gm IV q 6-8 hr
3	1 = no 2 = no 3 = no 4 = yes	<u>ampicillin 2 gm IV q 4h</u>  + ceftriaxone 2 gm IV q12 cefotaxime 2 gm IV q 6 h
4	1 = no 2 = no 3 = no 4 = no	<u>ceftriaxone 2 gm IV q 12 hr</u>  or cefotaxime 2 gm IV q 6 hr
5	1 = no 2 = no 3 = yes	<u>ampicillin 2 gm IV q 4 hr</u>  + ceftazidime 2 gm IV q 8 hr or cefipime 2 gm IV q 8 hr
6	1 = no 2 = yes 3 = no 4 = yes	<u>vancomycin 1.5 – 2 gm IV q 12 hr</u>  + chloramphenicol 1 gm IV q 6 hr
7	1 = no 2 = yes 3 = no 4 = no	
8	1 = no 2 = yes 3 = yes	
9	5 = yes to inputs 3-8	add to output consider acyclovir 10 mg/kg IV q 8h

In the Meningitis Input-Output Table, possible combinations of the five inputs are listed. For the conditions manifested in the patient, different drugs and dosages will be required. The proper treatment for each combination is listed in the output column of **Table x**. After the algorithm runs the comparison, the output is displayed on the computer screen, prompting the intensivist with the proper treatment **3712**.

Referring to **Figure 48**, the ventilator weaning decision support algorithm of the present invention is illustrated. The ventilator weaning decision support algorithm is used to determine whether an intensive care unit patient can return to breathing unassisted, and discontinue use of a

ventilator. Such a determination requires evaluation of the patient by the intensivist over the course of several days.

To begin the decision process of whether to wean a patient from ventilator use, the intensivist is prompted to conduct daily screening, preferably during the hours of 06:00 a.m. to 10:00 a.m. **3800**. The daily screen prompts the intensivist to determine whether: the patient's P/F ratio is greater than 200, the patient's positive end-expiratory pressure (PEEP) is less than or equal to 5, whether cough suctioning has been adequate and/or spontaneous, infusions with vasopressors have been necessary, and continuous infusions of sedatives or neuromuscular blocking agents have been necessary **3800**. If all conditions **3802** are answered no, the intensivist is directed by the system to repeat the daily screen **3805** the following morning. If all the conditions of the daily screen are met **3802**, the intensivist is prompted to perform additional tests.

If the patient has satisfied the daily screen, the intensivist is next directed to conduct a rapid shallow breathing test **3804**. To perform the test, the intensivist is directed to change the ventilator setting to continuous positive airway pressure (CPAP) less than or equal to 5. In other words, there is no intermittent mandatory ventilation or pressure support provided for the patient.

The patient is given one minute to reach a steady state of breathing. Then the intensivist measures the ratio of breaths per minute to tidal volume ( $f/V_T$ ). The intensivist next is prompted to determine whether the patient's  $f/V_T$  is less than or equal to 105 breathes per minute **3806**. If the patient's  $f/V_T$  is greater than 105 breathes per minute, the intensivist is prompted to return to performing daily screening the following morning **3808**.

If the patient's  $f/V_T$  is less than or equal to 105 breathes per minute, the intensivist is next

1 directed to perform a trial of spontaneous breathing. Here, the intensivist can either insert a T-  
2 Piece in the patient's airway or reduce the patient's CPAP to less than or equal to 5 over the  
3 course of two hours. The intensivist is prompted to observe the patient periodically in order to  
4 evaluate if the patient is breathing without assistance **3810**. The intensivist is prompted to  
5 perform a periodic assessment by determining whether: the patient's breathing characteristics  
6 are greater than 35 breaths per minute for 5 minutes, or SpO<sub>2</sub> is less than 90%, or the patient's  
7 Heart Rate (HR) is greater than 140, or HR deviates from the baseline breathing rate by more than  
8 20%, or the patient's SBP is outside the range of 90 to 180. If any of the conditions are met, the  
9 intensivist is directed by the system to terminate ventilator weaning **3812**. If the conditions are  
10 not met, the patient is further assessed.

11 In further assessment, the intensivist is prompted to determine whether the patient has  
12 been able to breathe spontaneously for two hours, keep a clear airway, and does not have any  
13 procedures scheduled within twenty-four hours that would require the patient to be intubated  
14 **3814**. If the patient meets all of these criteria **3814**, the intensivist is notified by the system that  
15 the patient may be extubated **3816**. If the patient does not meet one or more of the criteria **3814**,  
16 the intensivist is prompted to perform steps for progressive weaning **3818**.

17 Referring to **Figure 48A**, the ventilator weaning decision support algorithm of the  
18 present invention is further illustrated. The intensivist, at his or her discretion may choose  
19 either T-piece progressive weaning or pressure support progressive weaning. In order to perform  
20 T-piece progressive weaning, the intensivist is directed to repeat the trial of spontaneous  
21 breathing (as previously described **3810**). The intensivist can either insert a T-piece in the  
22 patient's airway or reduce the patient's CPAP to less than or equal to 5 over the course of two

hours. The intensivist is prompted to perform periodic assessment of the patient by either a two hour or 30 minute trial **3820**.

In order to perform pressure support progressive weaning, the intensivist is first prompted to observe whether the patient's pressure support (PS) rating is equal to eighteen plus or minus the positive end-expiratory pressure (PEEP). Next, the intensivist is directed by the system to regulate the pressure values in order to keep the patient's respiratory rate (RR) between twenty and thirty. Next, the intensivist is directed by the system to decrease the patient's pressure support by 2-4 centimeters of water two times per day. Once the patient maintains pressure support for at least two hours, the intensivist is prompted to further pursue extubating the patient **3822**.

After either T-Piece progressive weaning **3820** or pressure support progressive weaning **3822**, the intensivist is next prompted to perform a periodic assessment of the patient. Here, the intensivist must determine whether whether: the patient's breathing characteristics are greater than 35 breaths per minute for 5 minutes, or SpO<sub>2</sub> is less than 90%, or the patient's HR is grater than 140, or HR deviates from the baseline breathing rate by more than 20%, or the patient's SBP is outside the range of 90 to 180. Where the patient meets any of these criteria, the intensivist is prompted to terminate weaning. If the patient meets none of these criteria, the intensivist is prompted to further assess the patient's ability to breath spontaneously **3824**.

In further assessment, the intensivist is prompted to determine whether the patient has been able to breathe spontaneously for two hours, keep a clear airway, and does not have any procedures scheduled within twenty-four hours that would require the patient to be intubated **3826**. If the patient meets all of these criteria **3814**, the intensivist is notified by the system that



1 the patient may be extubated **3828**. If the patient does not meet one or more of the criteria **3826**,  
2 the intensivist is directed by the system to allow the patient to rest for at least twelve hours at  
3 A/C, the last level of pressure support the patient achieved **3830**. The intensivist is prompted to  
4 resume progressive weaning the following day **3832**.

5 Referring to **Figure 49**, the Warfarin Dosing Algorithm of the present invention is  
6 illustrated. The intensivist is first prompted to give the initial dose and determine subsequent  
7 dosage each day **3900**. When the intensivist determines subsequent dosage, he is first prompted  
8 to determine the patient's target INR **3902**. If the patient's target INR ranges from 2.0 to 3.0, the  
9 intensivist is prompted by the system to make further determinations relevant to dosage. The  
10 intensivist is directed by the system to determine whether the patient is taking drugs that effect  
11 prothrombin time **3904**, the baseline INR value **3906**, and whether rapid anticoagulation is  
12 required **3908**. Each answer is assigned a point value, and the total points are tabulated. If the  
13 point value is greater than one, the system refers to the 10 milligram load target database for  
14 dosing. If the point value is less than one, the system refers to the 5 milligram load target  
15 database for dosing **3910**.

16 At the initial INR determination **3902**, if the patient's INR was initially between 1.5 and  
17 2.0, the system refers to the 5 milligram load target database for dosing. If the patient's INR was  
18 initially between 3.0 and 4.0, the system refers to the 10 milligram load target database for  
19 dosing **3910**. Next the intensivist is prompted to enter the day of treatment **3912** and the  
20 patient's INR **3914**. Depending on whether the system has been directed to the 5 milligram load  
21 target or the 10 milligram load target, a comparison is run **3916** according to the following  
22 tables.

### 5 mg Load Target INR 1.5-2.0

day	<1.5	1.5-2	2-2.5	>2.5
2	5	1.25 - 2.5	0	0
3	5-7.5	1.25 - 2.5	0 - 1.25	0
4	10- (Check to see whether pt has received vit K)	1.25 - 2.5	0 - 1.25	0
5	10 (Check to see whether pt Has received vit K)	2.5 - 5	0 - 2.5	0 - 1.25
6	15 Obtain hematology consultation.	2.5 - 5	1.25 - 2.5	0 - 1.25

### 10 mg Load Target INR 3.0-4.0

day	<1.5	1.5-2	2-2.5	2.5-3	>3
2	10	7.5 - 10	5-7.5	2.5-5.0	0-2.5
3	10 -15	7.5 - 10	5-7.5	2.5 - 5	2.5-5
4	10 -15 (Check to see whether pt has received vit K)	7.5 -12.5	5 – 10	5-7.5	2.5-5
5	15 (Check to see whether pt has received vit K)	10 – 12.5	7.5-10	5 – 7.5	2.5-5
6	15-20 obtain hematology consultation.	10 - 15	7.5-12.5	5 - 10	5-7.5

The appropriate dosage and instructions is displayed on the computer screen to the intensivist 3918.

Referring to **Figure 51**, the heparin-induced thrombocytopenia (HIT) decision support algorithm of the present invention is illustrated. The intensivist is prompted to observe whether the patient's platelet count has dropped 50% or more over seventy-two hours while being treated with heparin, and whether any other obvious causes of platelet reduction might be present **4100**. If such a drop has not occurred, the intensivist is notified by the system that the patient most likely does not have HIT, but monitoring of the platelet count should continue **4102**. If the patient's platelet count has drastically dropped, the intensivist is prompted to determine whether the patient has been treated with heparin for more than three days **4104**. Regardless of the answer, the intensivist is next prompted to determine if the patient has been treated with heparin in the preceeding three months **4106**. If the patient has not received heparin in the preceeding three months, the intensivist is notified by the system that HIT is not likely to be the cause of the platelet drop. The intensivist is also prompted to monitor platelet count for infection or other thrombocytopenia-causing drugs, and to consider stopping heparin therapy if the platelet count drops below 50,000 per cubic millimeter **4108**.

If the patient has received heparin in the last three days **4104**, the intensivist is further prompted to look for signs of thrombosis, or blood clotting **4110**. If the patient shows signs of thrombosis, the intensivist is notified by the system that the patient is likely to have HIT. Accordingly, the intensivist is prompted to stop administering heparin and flush any drug administration equipment that would contain heparin traces. The intensivist is also provided instructions by the system to treat a patient still requiring anticoagulation treatment with alternate drugs and methods **4112**.

Where the patient does not show signs of thrombosis 4110, the intensivist is prompted to check for heparin resistance 4114. Signs of heparin resistance include inability to hold aPTT though heparin doses have been increase. If the patient shows signs of heparin resistance, the intensivist is prompted to consider stopping heparin treatment and to consider treating a patient still requiring anticoagulation treatment with alternate drugs and methods 4116. If the patient does not show signs of heparin resistance, the intensivist is notified by the system that the patient possibly has HIT. The intensivist is accordingly prompted to continue monitoring for thrombosis, consider infection or other drugs that cause thrombocytopenia, and to consider stopping heparin therapy if the platelet count drops below 50,000 per cubic millimeter 4118

## Results

The structure of the present invention and its efficacy have yielded striking results in practice. In a research setting, deployment of certain rudimentary aspects of the present the invention designed to experimentally test the approach described and developed in detail above, yielded unprecedented improvements in clinical and economic outcomes: 50% improvement in severity adjusted mortality, 40% improvement in clinical complication rates, 30% improvement in ICU length of stay, and 30% improvement in overall ICU cost of care.

A system and method of remote monitoring of ICU's and other healthcare locations has been shown. It will be apparent to those skilled in the art that other variations of the present invention are possible without departing from the scope of the invention as disclosed. For example, one can envision different ratios of command center/remote location to ICU's, other decision support algorithms that would be used by intensivists, other types of remote monitoring

1 of not only ICU's but other types of hospital functions as well as industrial functions where  
2 critical expertise is in limited supply but where that expertise must be applied to ongoing  
3 processes. In such cases a system such as that described can be employed to monitor processes  
4 and to provide standardized interventions across a number of geographically dispersed locations  
5 and operations.

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We claim:

1. A system for providing continuous, expert network health care services from a remote location comprising:
  - a plurality of health care locations;
  - at least one remote command center for managing healthcare at said plurality of health care locations; and at least one network;
  - wherein said plurality of health care locations are electronically connected to said at least one remote command center by the network, and wherein said at least one remote command center provides intensivist monitoring of the plurality of health care locations 24 hours per day, seven days per week.
2. The system for providing continuous, expert network health care services from a remote location of claim 1 wherein said remote command center further comprises a computerized patient care management system for monitoring and treating individual patients at any of said plurality of healthcare locations.
3. The system for providing continuous, expert network health care services from a remote location of claim 2 wherein said computerized patient care management system further comprises a data server/data warehouse for storing and analyzing data from the at least one remote command center.

4. The system for providing continuous, expert network health care services from a remote location of claim 1 wherein each of the plurality of health care locations further comprises patient monitoring equipment electronically connected to the at least one remote command center over the network.

5. The system for providing continuous, expert network health care services from a remote location of claim 4 wherein each health care location further comprises a nurses' station electronically connected to said monitoring equipment and to the at least one remote command center over the network.

6. The system for providing continuous, expert network health care services from a remote location of claim 1 wherein the healthcare locations comprise intensive care units (ICU's).

7. The system for providing continuous, expert network health care services from a remote location of claim 2 wherein said computerized patient care management system further comprises a relational database for storing a plurality of decision support algorithms and for prompting intensivists to provide care to patients based upon the any of the decision support algorithms.

8. The system for providing continuous, expert network health care services from a remote location of claim 7 wherein said algorithms are selected from the group consisting of algorithms for treating:

1 Acalculous Cholecystitis, Acute Pancreatitis Algorithm, Acute Renal Failure-Diagnosis, Acute  
2 Renal Failure-Management & Treatment, Adrenal Insufficiency, Agitation and Anxiety,  
3 Depression & Withdrawal, Aminoglycoside Dosing and Therapeutic Monitoring, an  
4 Amphotericin-B Treatment Guidelines, Analgesia, Antibiotic Classification & Costs,  
5 Antibiograms Algorithm, Antibiotic associated Colitis Algorithm, ARDS: Hemodynamic  
6 Management, ARDS: Steroid Use, ARDS: Ventilator Strategies, Asthma, Bleeding Patient,  
7 Bloodstream Infections, Blunt Cardiac Injury, Bradyarrhythmias, Brain Death, Bronchodilator  
8 Use in Ventilator Patients, Bronchoscopy & Thoracentesis Guidelines, Candiduria, Cardiogenic  
9 Shock, CardioPulmonary Resuscitation Guideline, Catheter Related Septicemia, a Catheter  
10 Replacement Strategies, Cervical Cord Injury, Congestive Heart Failure , Copd Exacerbation  
11 & Treatment, CXR (Indications), Dealing with Difficult patients and families, Diabetic  
12 Ketoacidosis, Dialysis, Diurectic Use, Drug Changes with Renal Dysfunction, Emergency  
13 Cardiac Pacing, Endocarditis Diagnosis and Treatment, Endocarditis Prophylaxis, End of Life  
14 Decisions, Endotracheal Tubes & Tracheotomy, Ethical Guidelines, Febrile Neutropenia, FUO,  
15 Fluid Resusditation, Guillain-Barre Syndrome, Heparin, Heparin-Induced Thrombocytopenia,  
16 Hepatic Encephalopathy, Hepatic Failure, HIV + Patent Infections, Hypercalcemia Diagnosis  
17 and Treatment, Hypercalcemia Insulin Treatment, Hyperkalemia : Etiology & Treatment,  
18 Hyponatremia : Etiology & Treatment, Hypertensive Crisis, Hypokalemia : Etiology &  
19 Treatment, Hyponatremia : Etiology & Treatment, Hypothermia, Identification of Cervical  
20 Cord Injury, Implantable Cardio-defibrillator, Intra-Aortic Balloon Device, Intracerebral  
21 Hemorrhage, Latex Allergy, Magnesium Administration, Management of Hypotension,  
22 Inotropes , Management of Patients with Ascites, Empiric Meningitis, Meningitis,a Myasthenia



Gravis, Myocardial Infarction, Myocardial Infarction with left bundle branch block, Necrotizing Soft Tissue Infections, Neuromuscular Blockers, Neuromuscular Complications of Critical Illness, Non-Infectious Causes of Fever, Non-Traumatic Coma, Noninvasive Modes of Ventilation, Nutritional Management, Obstetrical Complication, Oliguria, Oliguria, Open Fractures, Open Fractures, Ophthalmic Infections, Organ Procurement Guidelines, PA Catheter Guideline and Troubleshooting, Pancreatitis, Penetrating Abdominal Injury, Penetrating Chest Injury, Penicillin Allergy, Penicillin Allergy, Permanent Pacemaker and Indications, Pneumonia Community Acquired, Pneumonia Hospital Acquired, Post-Op Bleeding, Post-Op Hypertension, Post-Op Hypertension , Post-Op Management of Abdominal, Post-Op Management of Carotid, Post-Op Management of Open Heart, Post-Op Management of Thoracotomy, Post-Op Myocardial Ischemia (Non-Cardiac Arrhythmias after Cardiac Surgery), Post-Op Power Weaning, Pressure Ulcers, Pulmonary Embolism Diagnosis, Pulmonary Embolism Treatment, Respiratory Isolation, Sedation, Seizure, Status Epilepticus, Stroke, Sub-Arachnoid Hemorrhage, Supra-Ventricular Tachyarrhythmia, Supra-Ventricular Tachycardia, Wide Complex QRS Tachycardia, Therapeutic Drug Monitoring, Thrombocytopenia, Thrombolytic Therapy, Transfusion Guidelines, Traumatic Brain Injury, Assessment of Sedation, Sedation, Septic Shock, Bolus Sliding Scale Midazolam, Short Term Sedation Process, Sinusitis, SIRS, Spinal Cord Injury, Steroid Replacement Strategy, Thyroid Disease, Transplant Infection Prophylaxis, Transplant Related Infections, Treatment of Airway Obstruction, Unknown Poisoning, Unstable Angina, Upper GI Bleeding Stress Prophylaxis, Vancomycin, Upper GI Bleeding Non-Variceal, Upper GI Bleeding Variceal , Use of Hematopoietic Growth Factors, Ventilation Weaning, Ventilation Weaning Protocol, Venous

Thrombosis Diagnostic and Treatment, Venous Thromboembolism Prophylaxis, Ventricular Arrhythmia, Warfarin, Warfarin Dosing, and Wound Healing Strategies.

10. The system for providing continuous, expert network health care services from a remote location of claim 2 wherein said computerized patient care management system further comprises order writing software for providing knowledge-based recommendations and prescriptions for medication based upon the clinical data.

11. The system for providing continuous, expert network health care services from a remote location of claim 2 wherein said computerized patient care management system further comprises knowledge-based vital sign/hemodynamic algorithms that prompt said intensivist to engage in early intervention.

12. A method for providing continuous expert critical care comprising:  
monitoring patients in a plurality of ICU's;  
communicating the information from the patient monitoring to at least one command center over a first network  
receiving and analyzing the information from the patient monitoring at the command center over the first network; and  
providing guidance from the command center to the plurality of ICU's to take actions regarding patient care.

13. The method for providing continuous expert critical care of claim 12 wherein the providing guidance from the command center further comprises an intensivist reviewing decision support algorithms that provide guidance for treating a plurality of critical care conditions.

14. The method for providing continuous expert critical care of claim 13 wherein the decision support algorithms are taken from the group consisting of algorithms for treating: Acalculous Cholecystitis, Acute Pancreatitis Algorithm, Acute Renal Failure-Diagnosis, Acute Renal Failure-Management & Treatment, Adrenal Insufficiency, Agitation and Anxiety, Depression & Withdrawal, Aminoglycoside Dosing and Therapeutic Monitoring, an Amphotericin-B Treatment Guidelines, Analgesia, Antibiotic Classification & Costs, Antibigrams Algorithm, Antibiotic associated Colitis Algorithm, ARDS: Hemodynamic Management, ARDS: Steroid Use, ARDS: Ventilator Strategies, Asthma, Bleeding Patient, Bloodstream Infections, Blunt Cardiac Injury, Bradyarrhythmias, Brain Death, Bronchodilator Use in Ventilator Patients, Bronchoscopy & Thoracentesis Guidelines, Candiduria, Cardiogenic Shock, CardioPulmonary Resuscitation Guideline, Catheter Related Septicemia, a Catheter Replacement Strategies, Cervical Cord Injury, Congestive Heart Failure , Copd Exacerbation & Treatment, CXR (Indications), Dealing with Difficult patients and families, Diabetic Ketoacidosis, Dialysis, Diuretic Use, Drug Changes with Renal Dysfunction, Emergency Cardiac Pacing, Endocarditis Diagnosis and Treatment, Endocarditis Prophylaxis, End of Life Decisions, Endotracheal Tubes & Tracheotomy, Ethical Guidelines, Febrile Neutropenia, FUO, Fluid Resusditation, Guillain-Barre Syndrome, Heparin, Heparin-Induced Thrombocytopenia, Hepatic Encephalopathy, Hepatic Failure, HIV + Patent Infections, Hypercalcemia Diagnosis

and Treatment, Hypercalcemia Insulin Treatment, Hyperkalemia : Etiology & Treatment, Hybern timers  
Hypernatremia : Etiology & Treatment, Hypertensive Crisis, Hypokalemia : Etiology & Treatment, Hyponatremia : Etiology & Treatment, Hypothermia, Identification of Cervical Cord Injury, Implantable Cardio-defibrillator, Intra-Aortic Balloon Device, Intracerebral Hemorrhage, Latex Allergy, Magnesium Administration, Management of Hypotension, Inotropes , Management of Patients with Ascites, Empiric Meningitis, Meningitis,a Myasthenia Gravis, Myocardial Infarction, Myocardial Infarction with left bundle branch block, Necrotizing Soft Tissue Infections, Neuromuscular Blockers, Neuromuscular Complications of Critical Illness, Non-Infectious Causes of Fever, Non-Traumatic Coma, Noninvasive Modes of Ventilation, Nutritional Management, Obstetrical Complication, Oliguria, Oliguria, Open Fractures, Open Fractures, Ophthalmic Infections, Organ Procurement Guidelines, PA Catheter Guideline and Troubleshooting, Pancreatitis, Penetrating Abdominal Injury, Penetrating Chest Injury, Penicillin Allergy, Penicillin Allergy, Permanent Pacemaker and Indications, Pneumonia Community Acquired, Pneumonia Hospital Acquired, Post-Op Bleeding, Post-Op Hypertension, Post-Op Hypertension , Post-Op Management of Abdominal, Post-Op Management of Carotid, Post-Op Management of Open Heart, Post-Op Management of Thoracotomy, Post-Op Myocardial Ischemia (Non-Cardiac Arrhythmias after Cardiac Surgery), Post-Op Power Weaning, Pressure Ulcers, Pulmonary Embolism Diagnosis, Pulmonary Embolism Treatment, Respiratory Isolation, Sedation, Seizure, Status Epilepticus, Stroke, Sub-Arachnoid Hemorrhage, Supra-Ventricular Tachyarrhythmia, Supra-Ventricular Tachycardia, Wide Complex QRS Tachycardia, Therapeutic Drug Monitoring, Thrombocytopenia, Thrombolytic Therapy, Transfusion Guidelines, Traumatic Brain Injury,

1    Assessment of Sedation, Sedation, Septic Shock, Bolus Sliding Scale Midazolam, Short Term  
2    Sedation Process, Sinusitis, SIRS, Spinal Cord Injury, Steroid Replacement Strategy, Thyroid  
3    Disease, Transplant Infection Prophylaxis, Transplant Related Infections, Treatment of Airway  
4    Obstruction, Unknown Poisoning, Unstable Angina, Upper GI Bleeding Stress Prophylaxis,  
5    Vancomycin, Upper GI Bleeding Non-Variceal, Upper GI Bleeding Variceal , Use of  
6    Hematopoietic Growth Factors, Ventilator Weaning, Ventilator Weaning Protocol, Venous  
7    Thrombosis Diagnostic and Treatment, Venous Thromboembolism Prophylaxis, Ventricular  
8    Arrhythmia, Warfarin, Warfarin Dosing, and Wound Healing Strategies.

9  
10   14. The method for providing continuous expert critical care of claim 13 further comprising a  
11   data server/ data warehouse storing and analyzing patient data from the at least one command  
12   center and providing analysis in results over a second network to the at least one command  
13   center.

1           **Abstract of the Disclosure**

2  
3           A system and method for providing continuous expert network critical care services from  
4 a remote location. A plurality of intensive care units (ICU's) with associated patient monitoring  
5 instrumentation is connected over a network to a command center which is manned by  
6 intensivists 24 hours a day, 7 days a week. The intensivists are prompted to provide critical care  
7 by a standardized series of guideline algorithms for treating a variety of critical care conditions.  
8 Intensivists monitor the progress of individual patients at remote intensive care units. A smart  
9 alarm system provides alarms to the intensivists to alert the intensivists to potential patient  
10 problems so that intervention can occur in a timely fashion. A data storage/data warehouse  
11 function analyzes individual patient information from a plurality of command centers and  
12 provides updated algorithms and critical care support to the command centers.

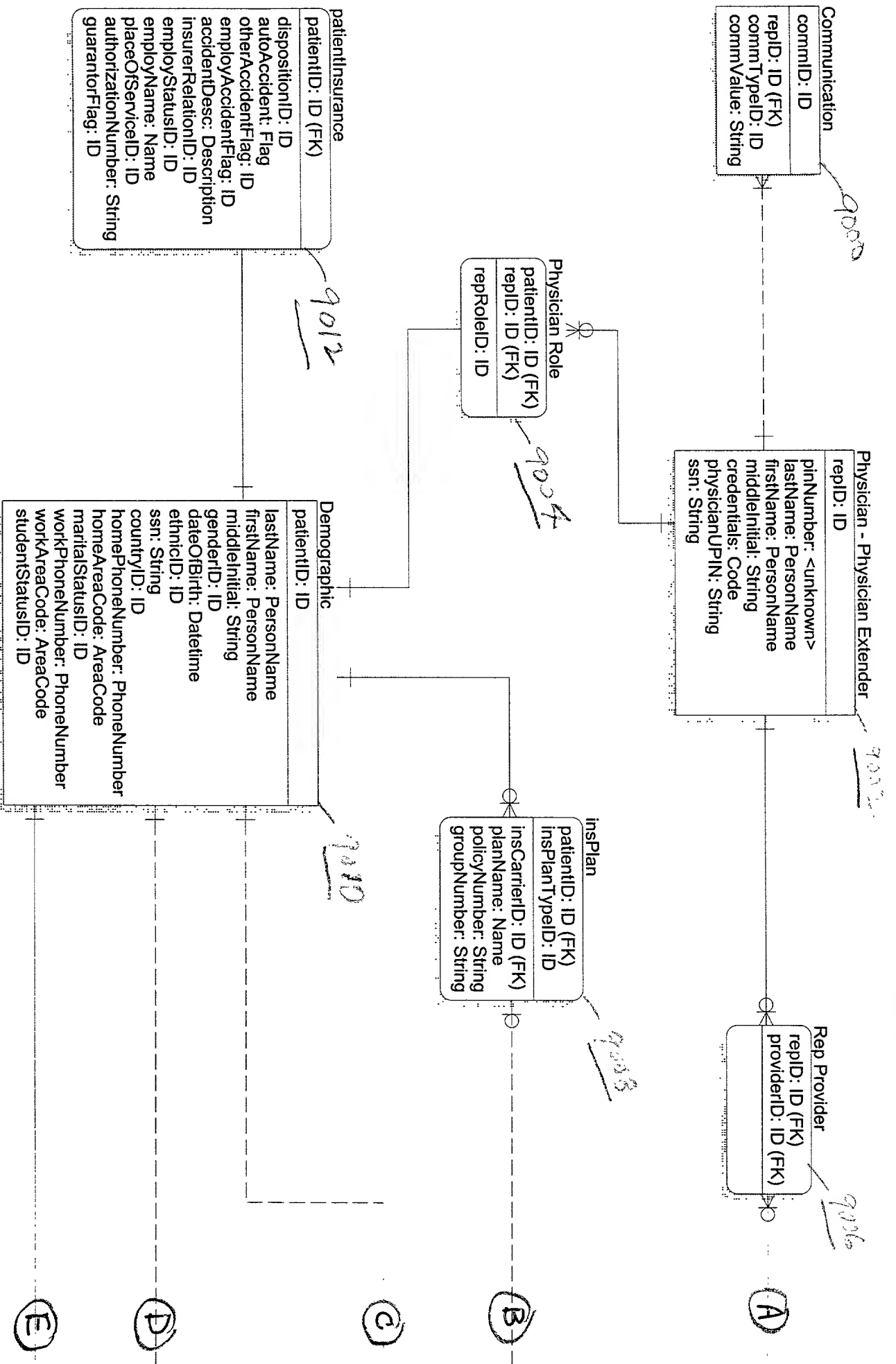


Fig. 1

(A)

Provider Number
providerID: ID
physicianProviderNumber: String
providerTypeID: ID

9014

(B)

Insurance Carrier
insCarrierID: ID
insCarrierName: Name
addressLine1: Address
addressLine2: Address
city: Name
stateID: ID
zipCode: ZipCode
zipCodeExtension: ZipCode

9018

(B)

Patient Visit
visitID: ID
patientID: ID (FK)
hospDischargeFlag: Flag
icuDischargeFlag: Flag
dateEntered: Datetime
patientStatusID: ID
icuAdmitDate: Datetime
hospAdmitDate: Datetime
medicalRecordNumber: String
sourceID: ID
hospDischargeDate: Datetime
hospDischargeID: ID
icuDischargeDate: Datetime
icuDischargeID: ID
readmitFlag: ID

9017

(C)

patientAddress
patientID: ID (FK)
addressID: ID (FK)

9020

(D)

Address
addressID: ID
addressTypeID: ID
addressLine1: Address
addressLine2: Address
city: Name
stateID: ID
zipCode: ZipCode
zipCodeExtension: ZipCode

9022

(E)

Guarantor
guarantorID: ID
patientID: ID (FK)
lastName: PersonName
firstName: PersonName
middleInitial: String
genderID: ID
dateOfBirth: Datetime
raceID: ID
ssn: String
countryID: ID
homePhoneNumber: PhoneNumber
homeAreaCode: Number
maritalStatusID: ID
workPhoneNumber: PhoneNumber
workAreaCode: Number
studentStatusID: ID

9026

guarantorAddress
guarantorID: ID (FK)
addressID: ID (FK)

9024



Fig. 2

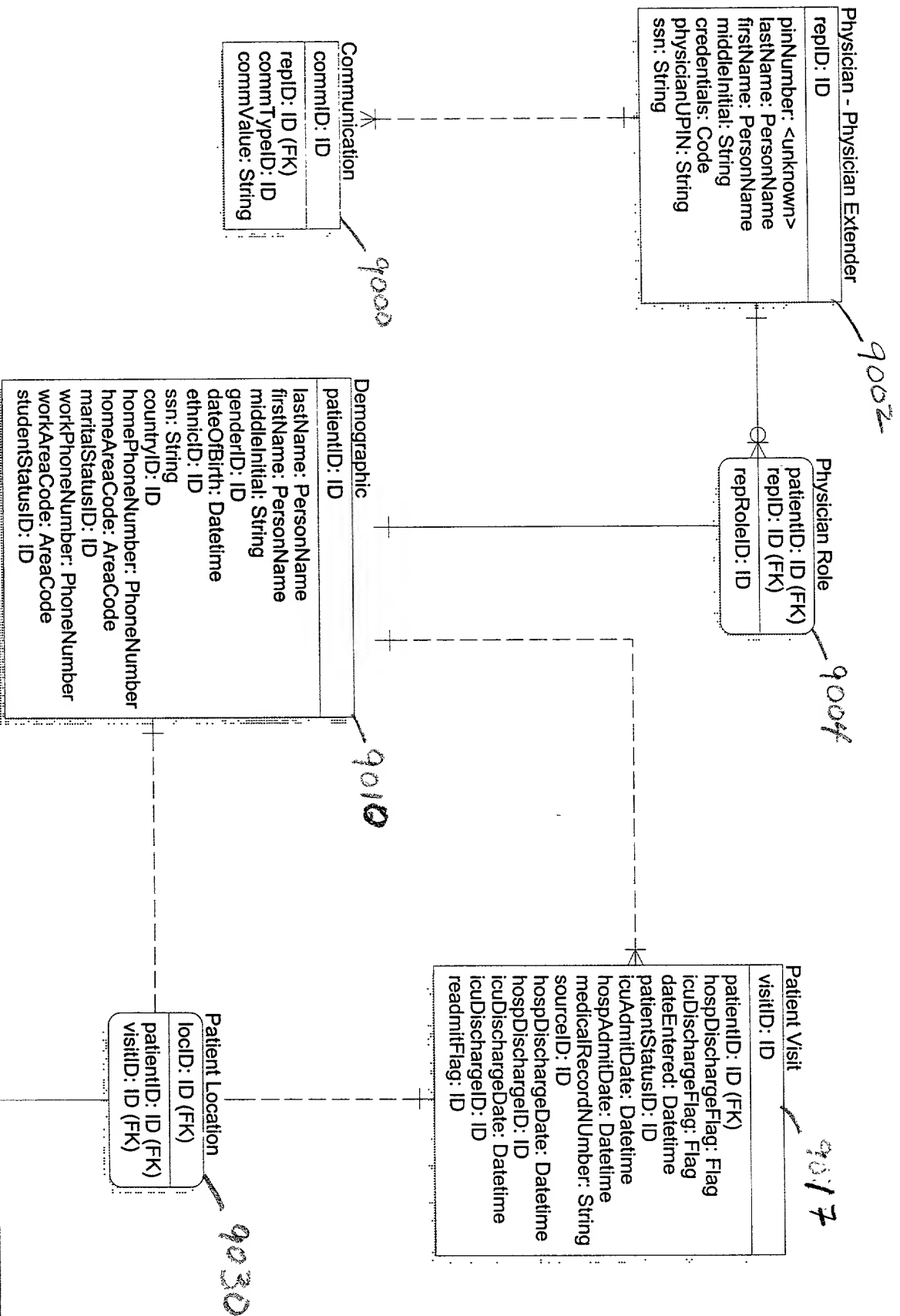


FIG. 2A

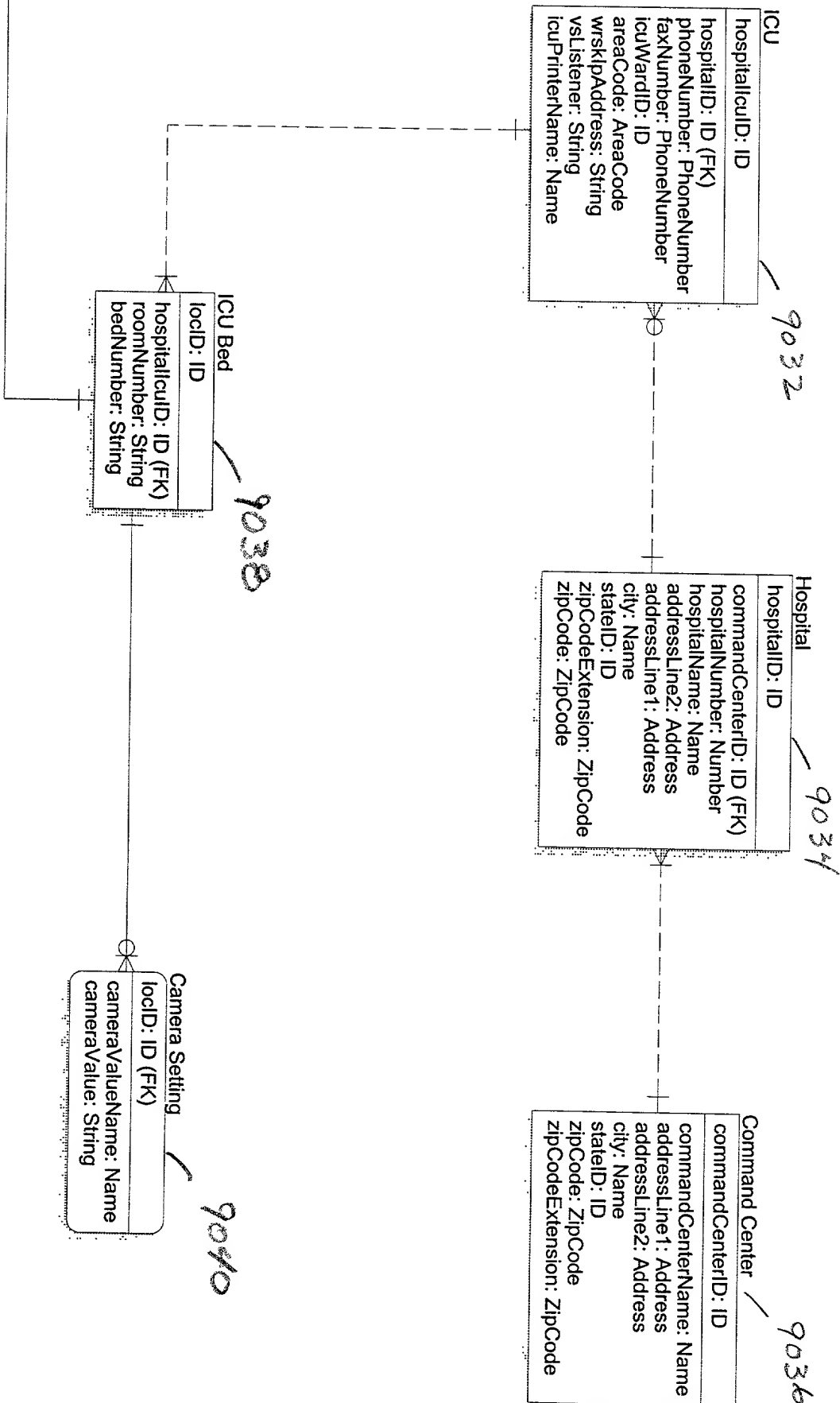


Fig. 3

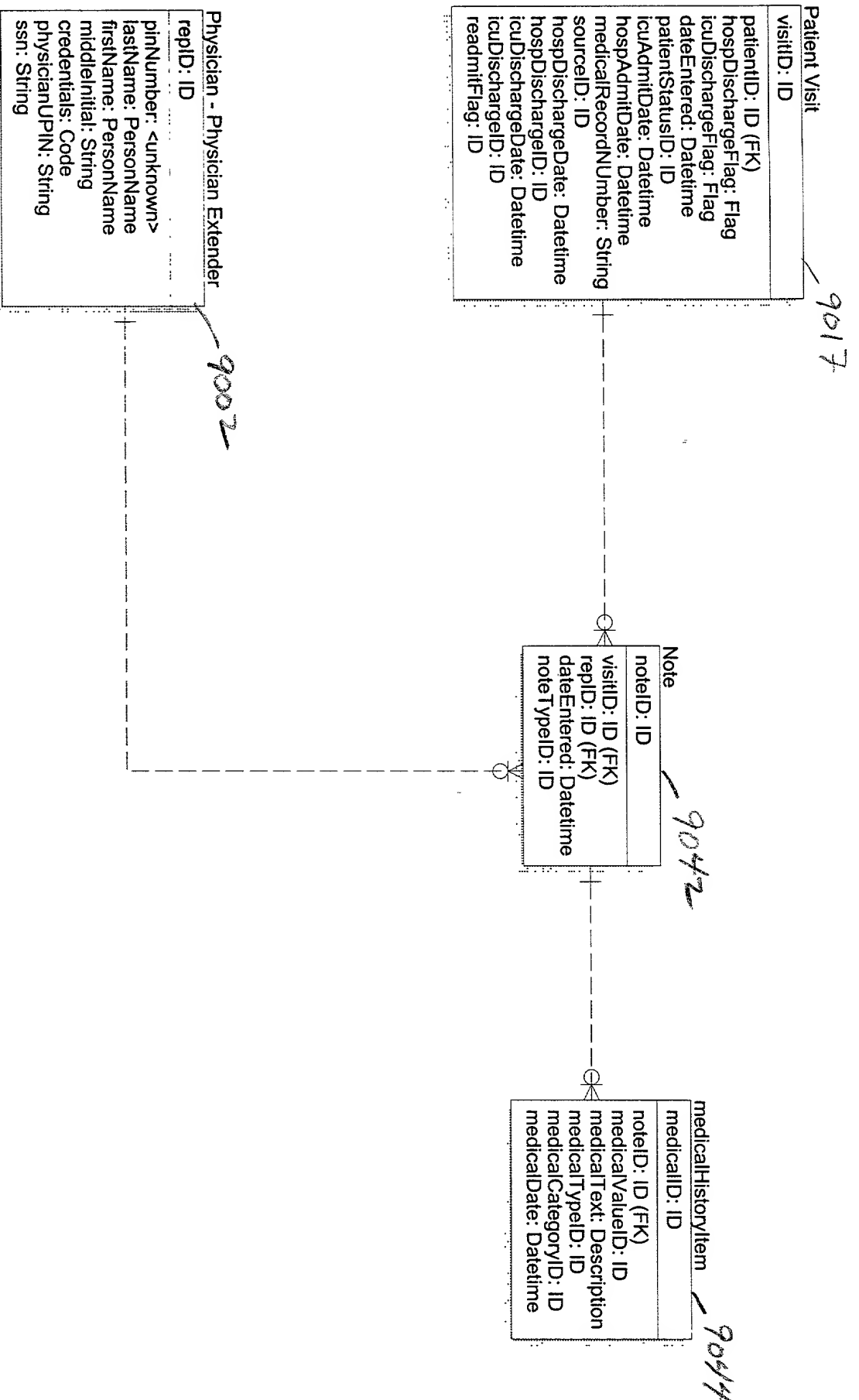


Fig. 4

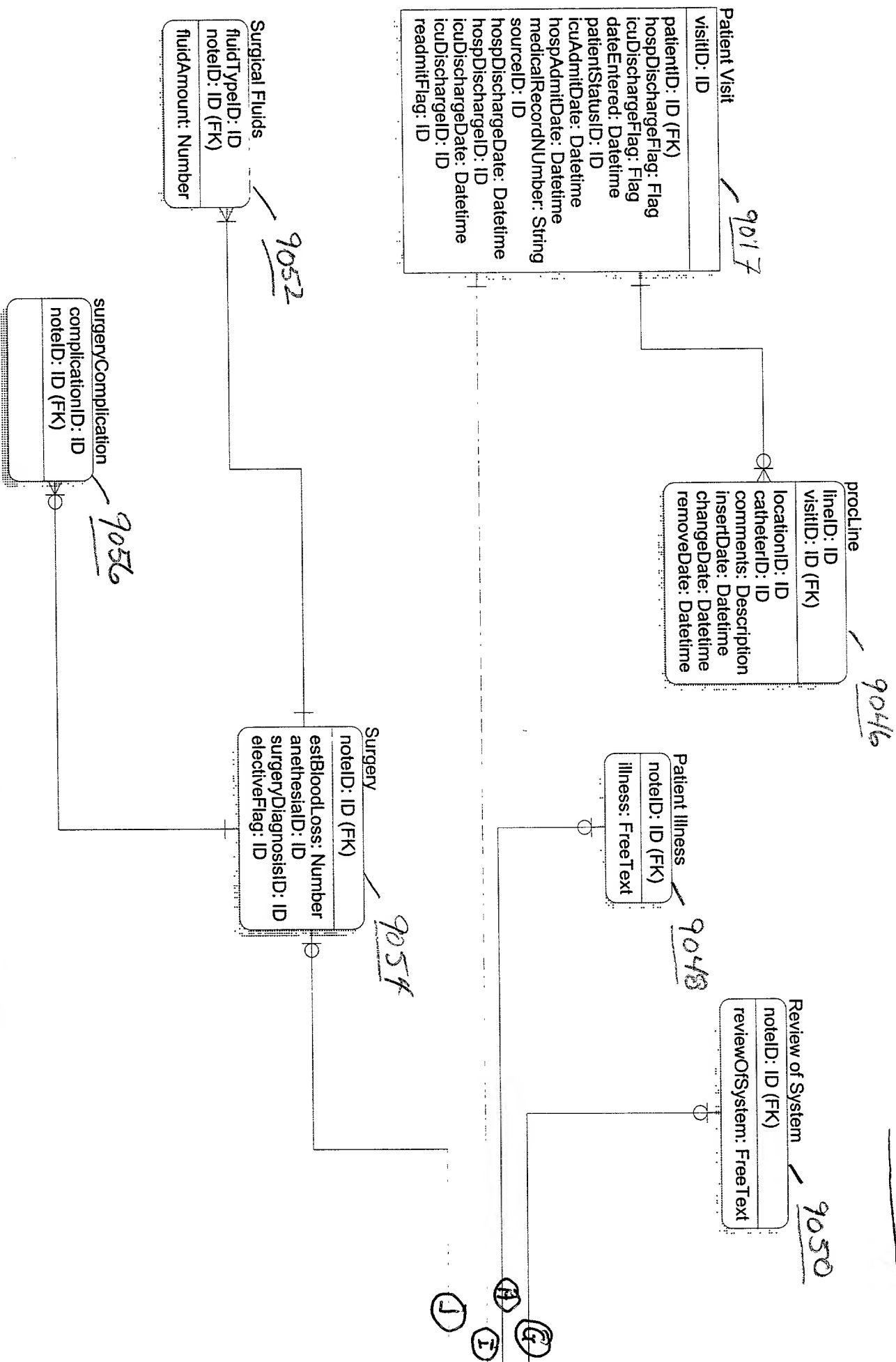




Fig. 4B

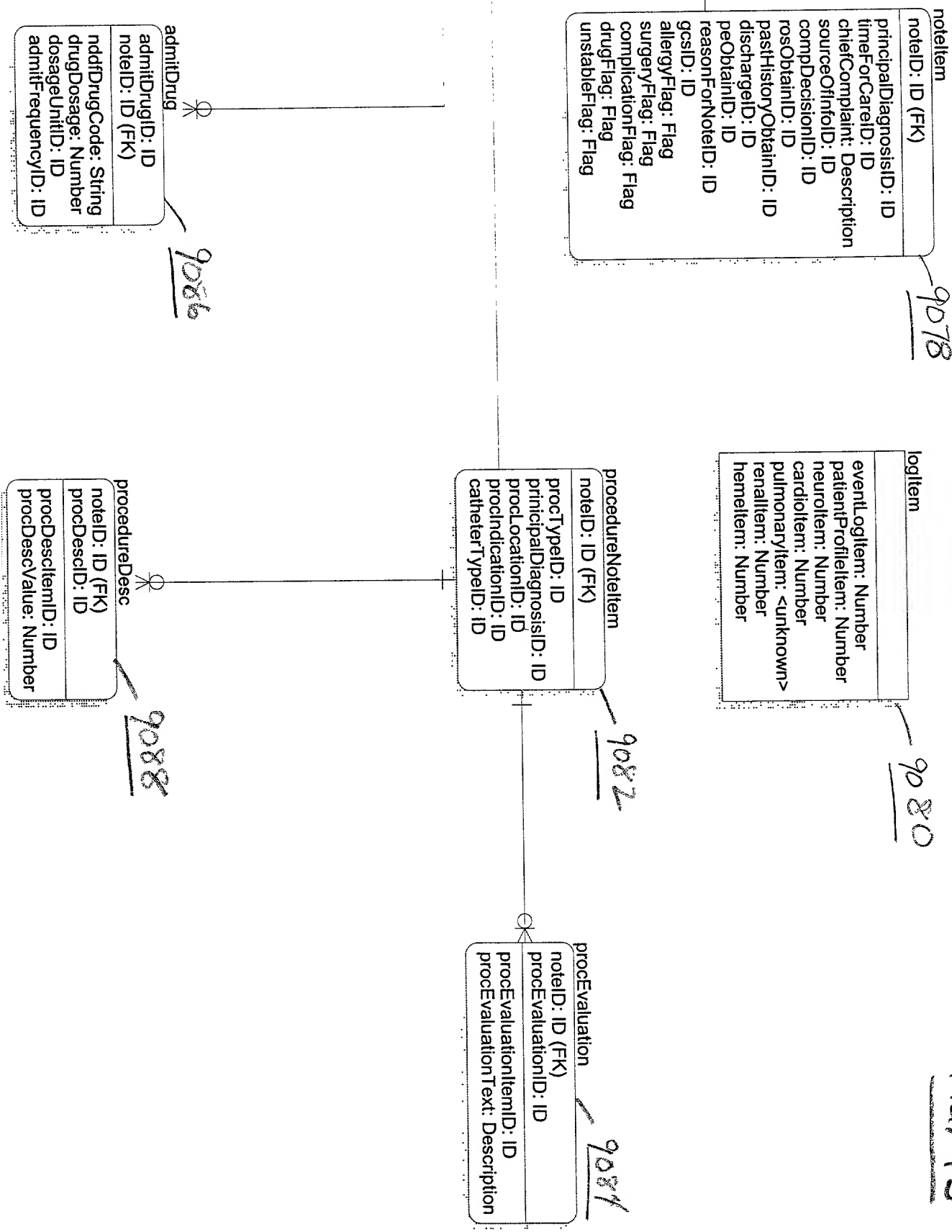
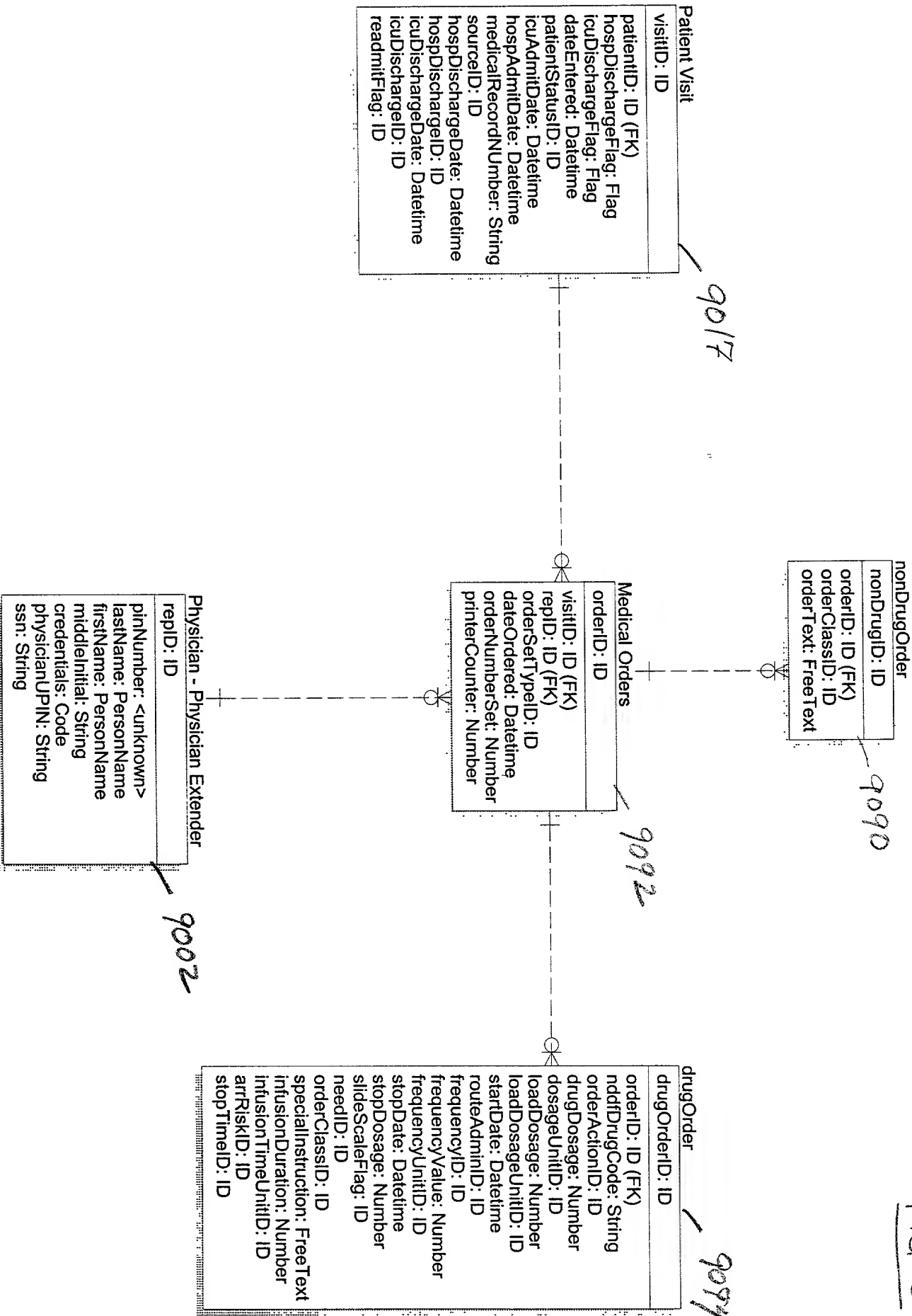
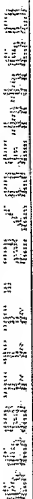


Fig 5







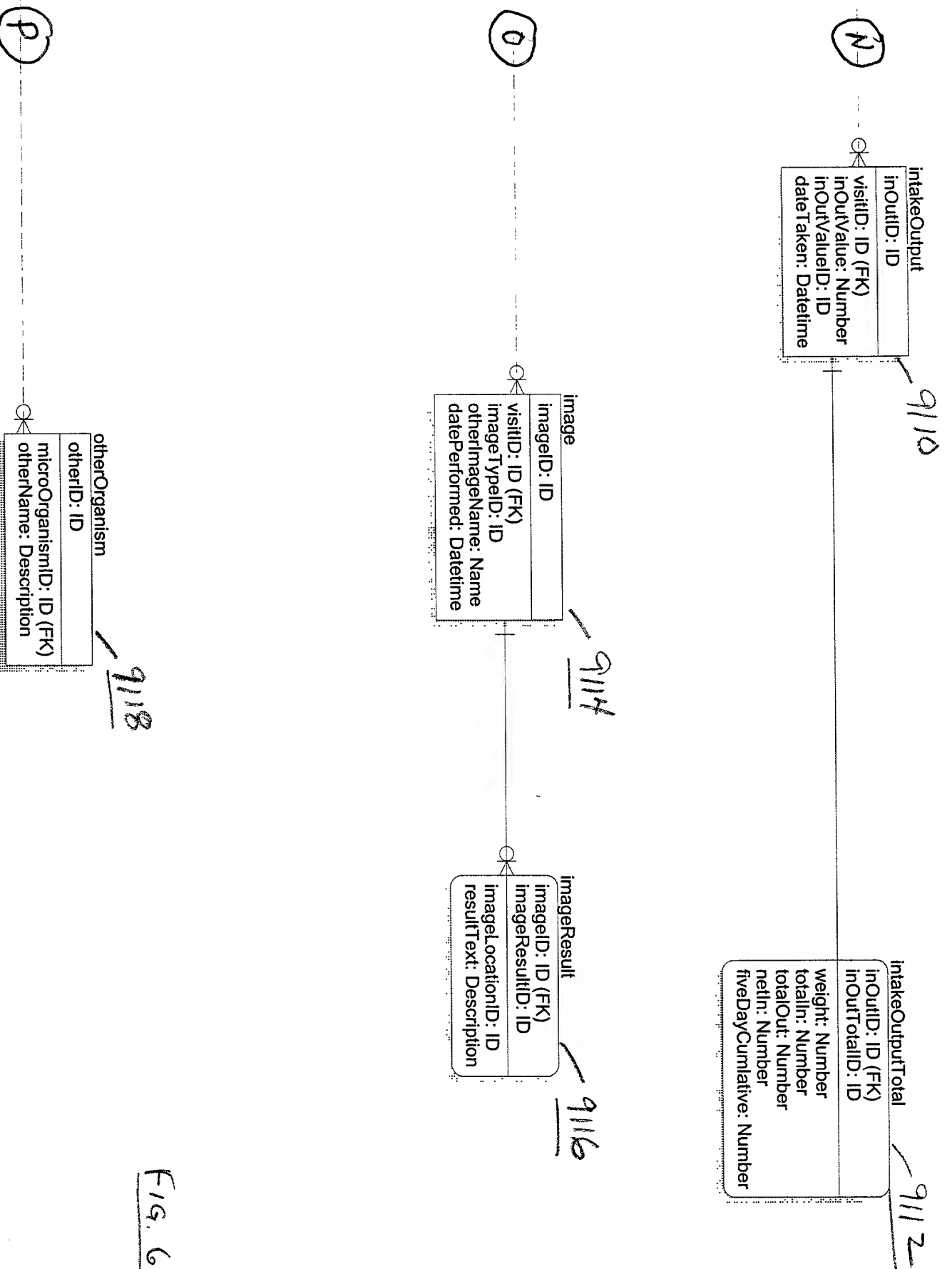


Fig. 6A

Fig 7

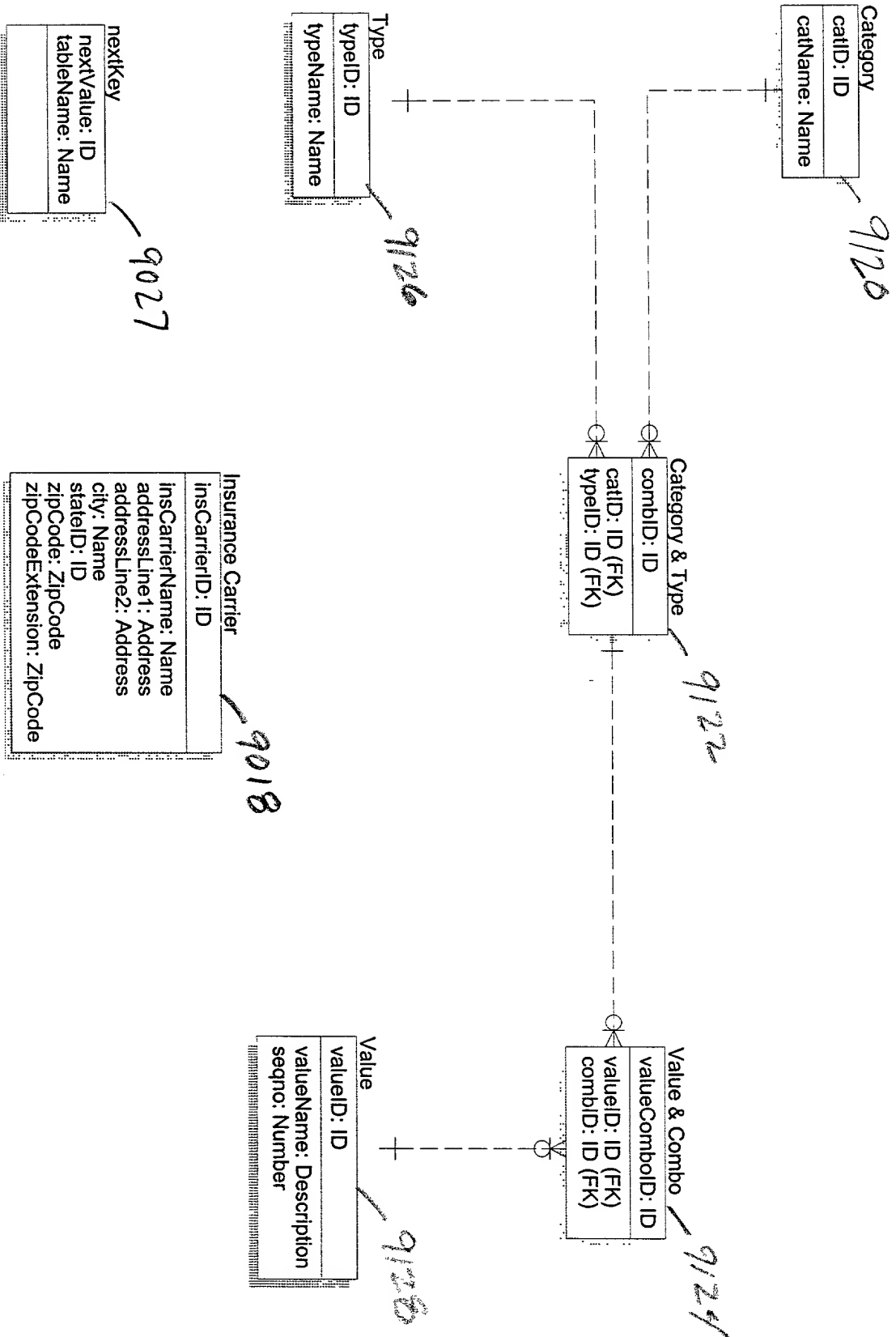


FIG 8

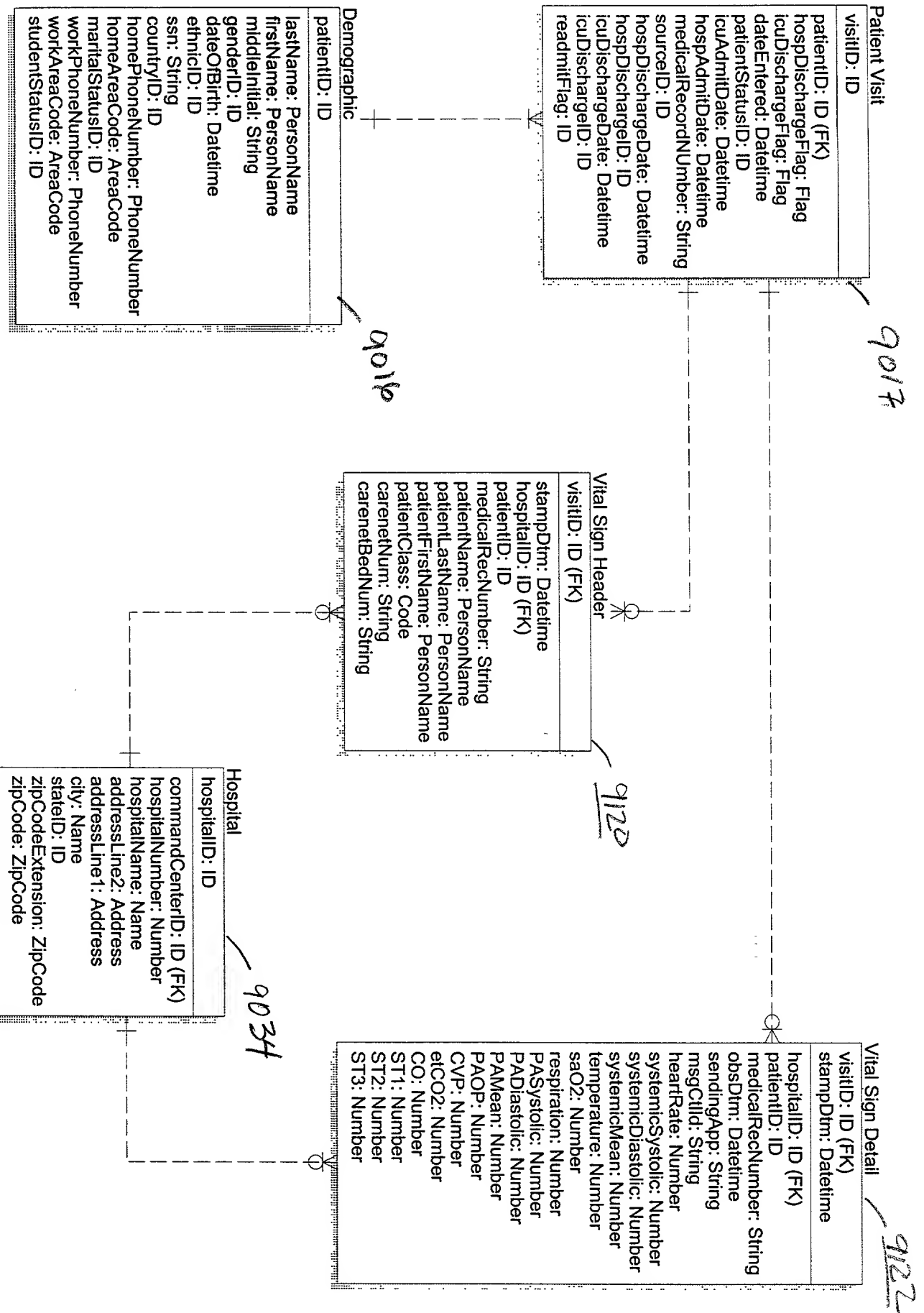


Fig. 8A

Vital Sign Log Detail	
hospitalID: ID (FK)	medicalRecNum: String (FK)
stampDtm: Datetime	
obsDtm: Datetime	
sendingApp: String	
msgCtId: String	
heartRate: Number	
systemicSystolic: Number	
systemicDiastolic: Number	
systemicMean: Number	
temperature: Number	
saO2: Number	
respiration: Number	
PASystolic: Number	
PADiastolic: Number	
PAMean: Number	
PAOP: Number	
CVP: Number	
etCO2: Number	
CO: Number	
ST1: Number	
ST2: Number	
ST3: Number	

9124

Vital Sign Log Header	
hospitalID: ID	medicalRecNum: String
stampDtm: Datetime	
patientName: PersonName	
patientLastName: PersonName	
patientFirstName: PersonName	
patientClass: Code	
carenetNum: String	
carenetBedNum: String	

9120

CaretPatientLocation	
hospitalID: ID	carenetNum: String
	carenetBedNum: String
locID: ID (FK)	

9132

ICU Bed	
locID: ID	hospitalCulID: ID (FK)
	roomNum: String
	bedNum: String

9038

Vital Sign Error Detail	
hospitalID: ID (FK)	medicalRecNum: String (FK)
stampDtm: Datetime	
code: String (FK)	
obsDtm: Datetime	
sendingApp: String	
msgCtId: String	
heartRate: Number	
systemicSystolic: Number	
systemicDiastolic: Number	
systemicMean: Number	
temperature: Number	
saO2: Number	
respiration: Number	
PASystolic: Number	
PADiastolic: Number	
PAMean: Number	
PAOP: Number	
CVP: Number	
etCO2: Number	
CO: Number	
ST1: Number	
ST2: Number	
ST3: Number	

9126

Vital Sign Error Header	
hospitalID: ID	medicalRecNum: String
stampDtm: Datetime	
patientName: PersonName	
patientLastName: String	
patientFirstName: String	
patientClass: Code	
carenetNum: String	
carenetBedNum: String	

9130

Vital Sign ErrorCode	
code: String	
description: String	

9118

# Distributed Architecture

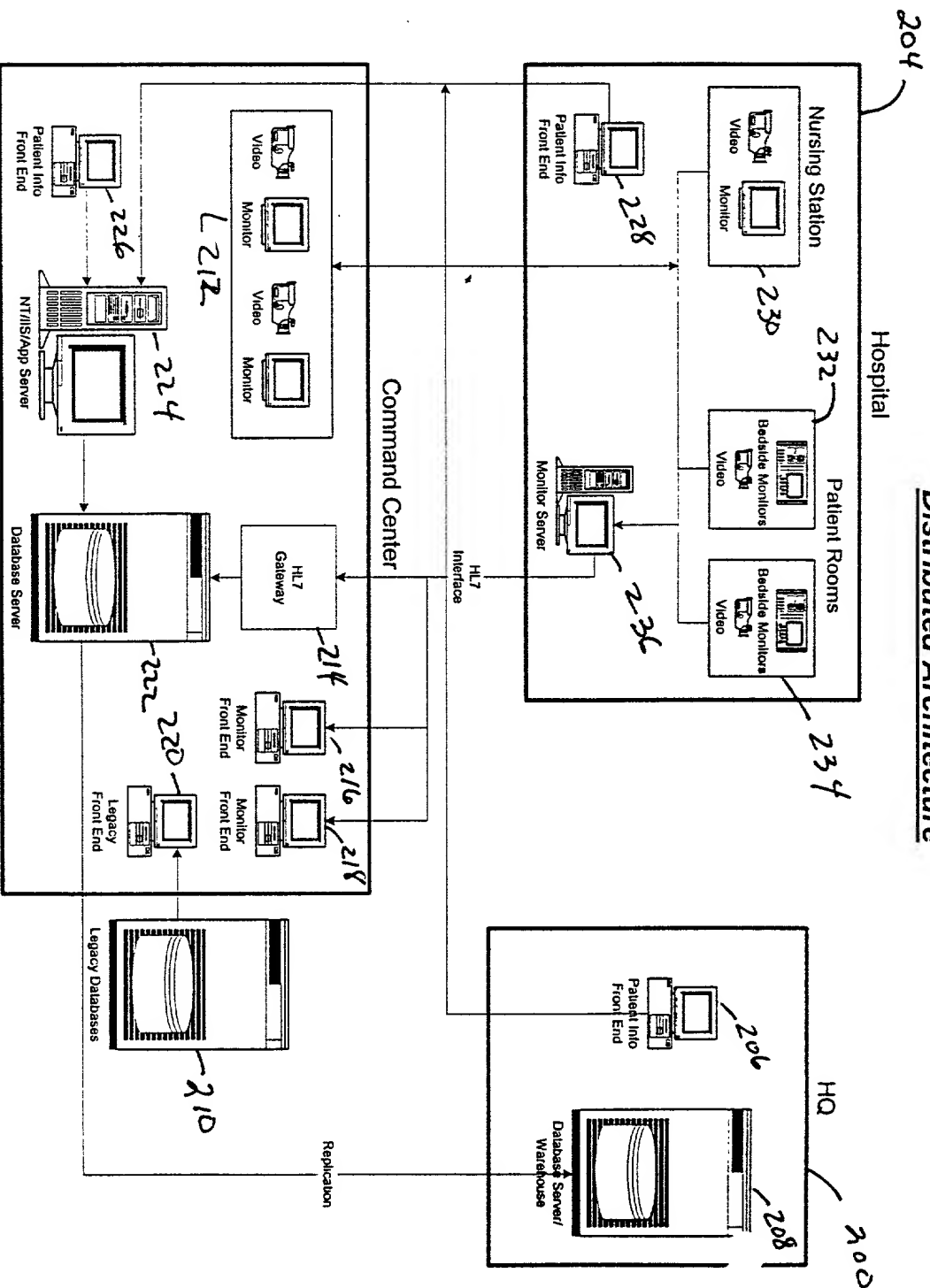
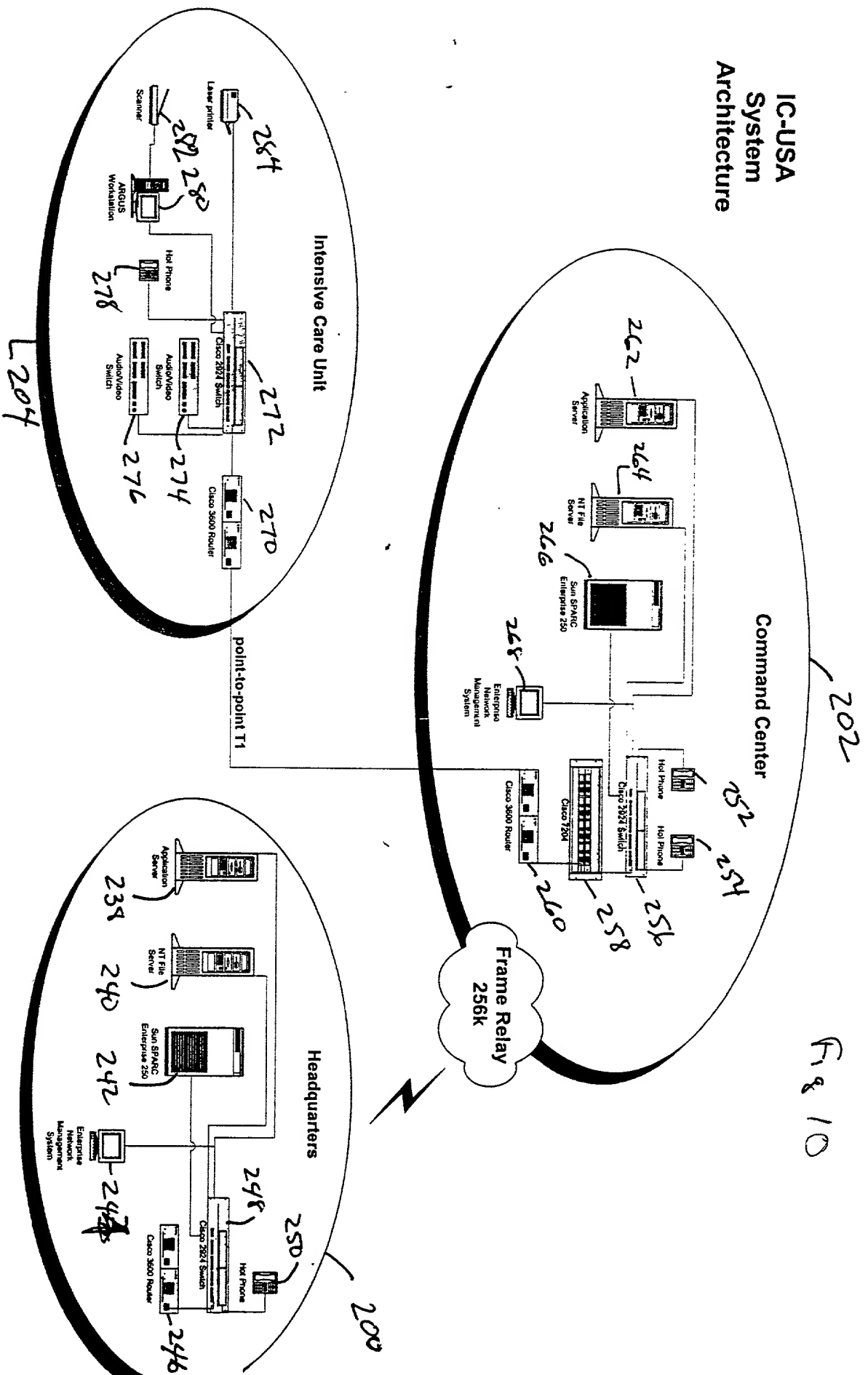
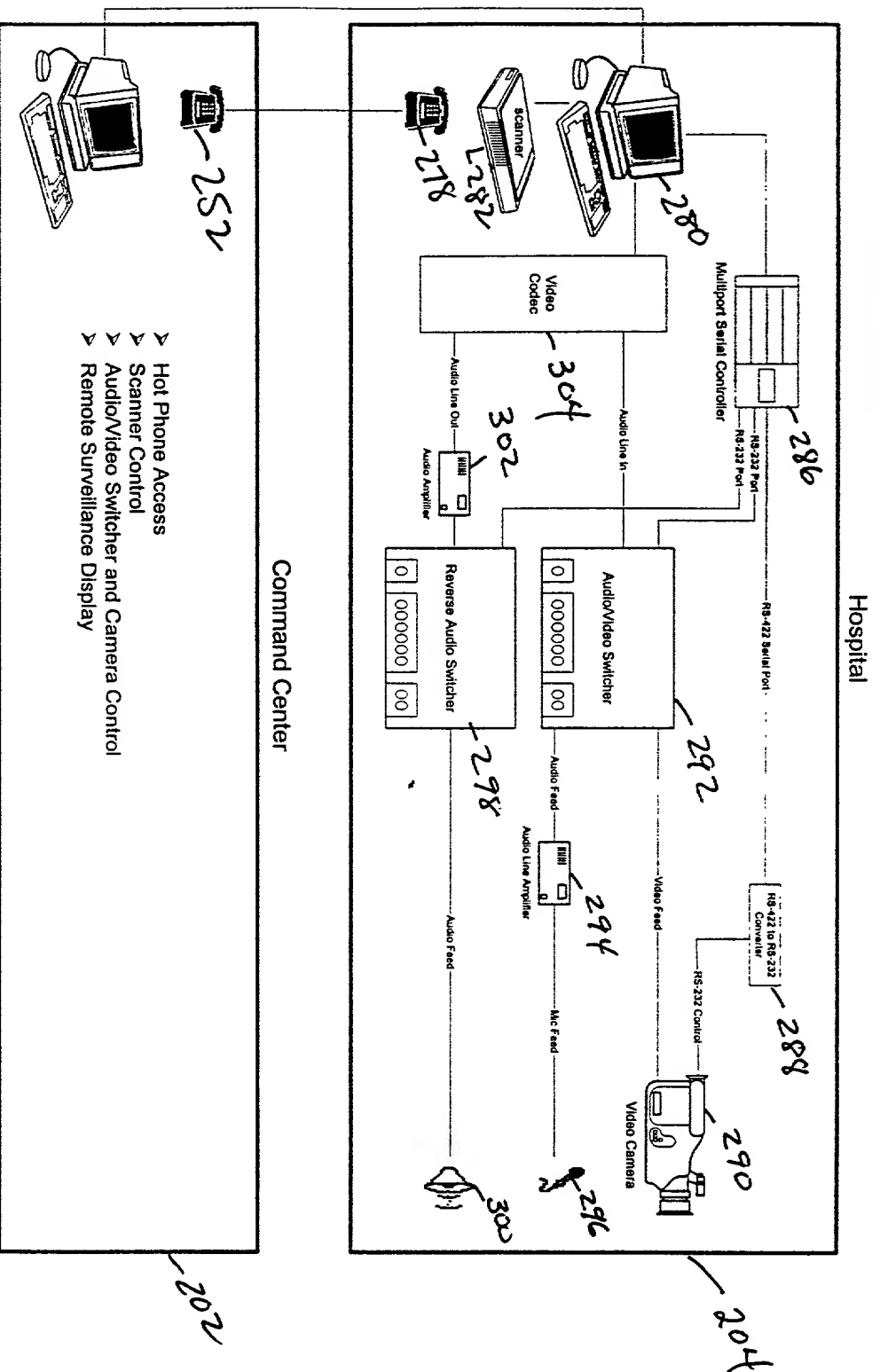


Fig. 9

# IC-USA System Architecture



# Video Conferencing/Surveillance/Imaging Components



# Vital Signs Data Flow

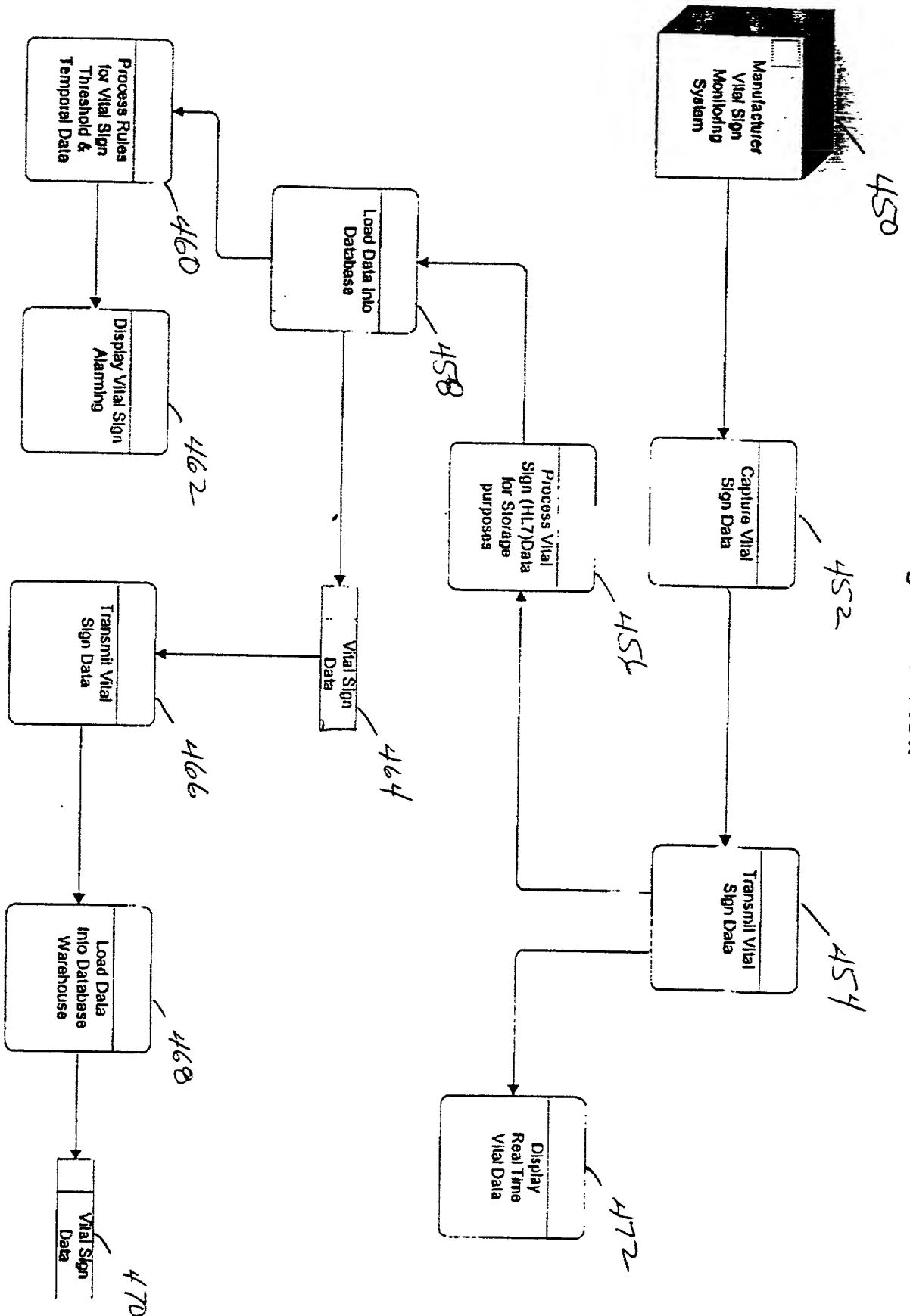


Fig 12

09443072 111899



# Patient Interaction Data Flow

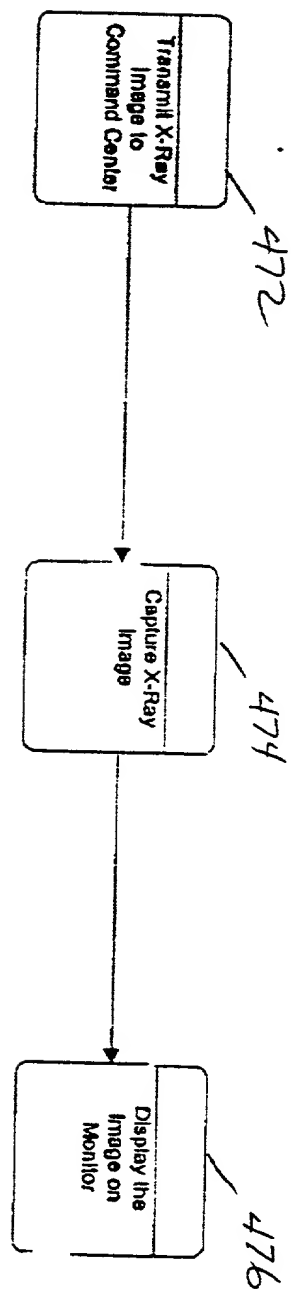


Fig 13A

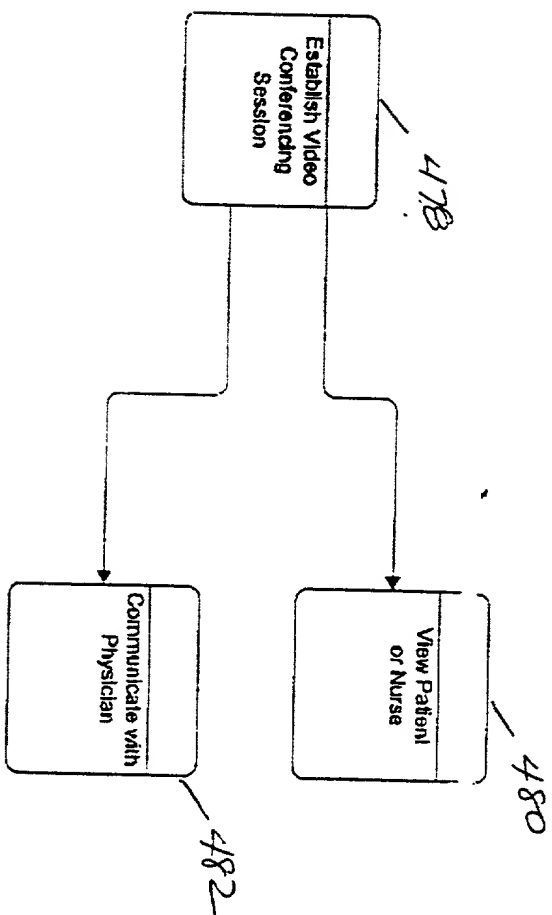
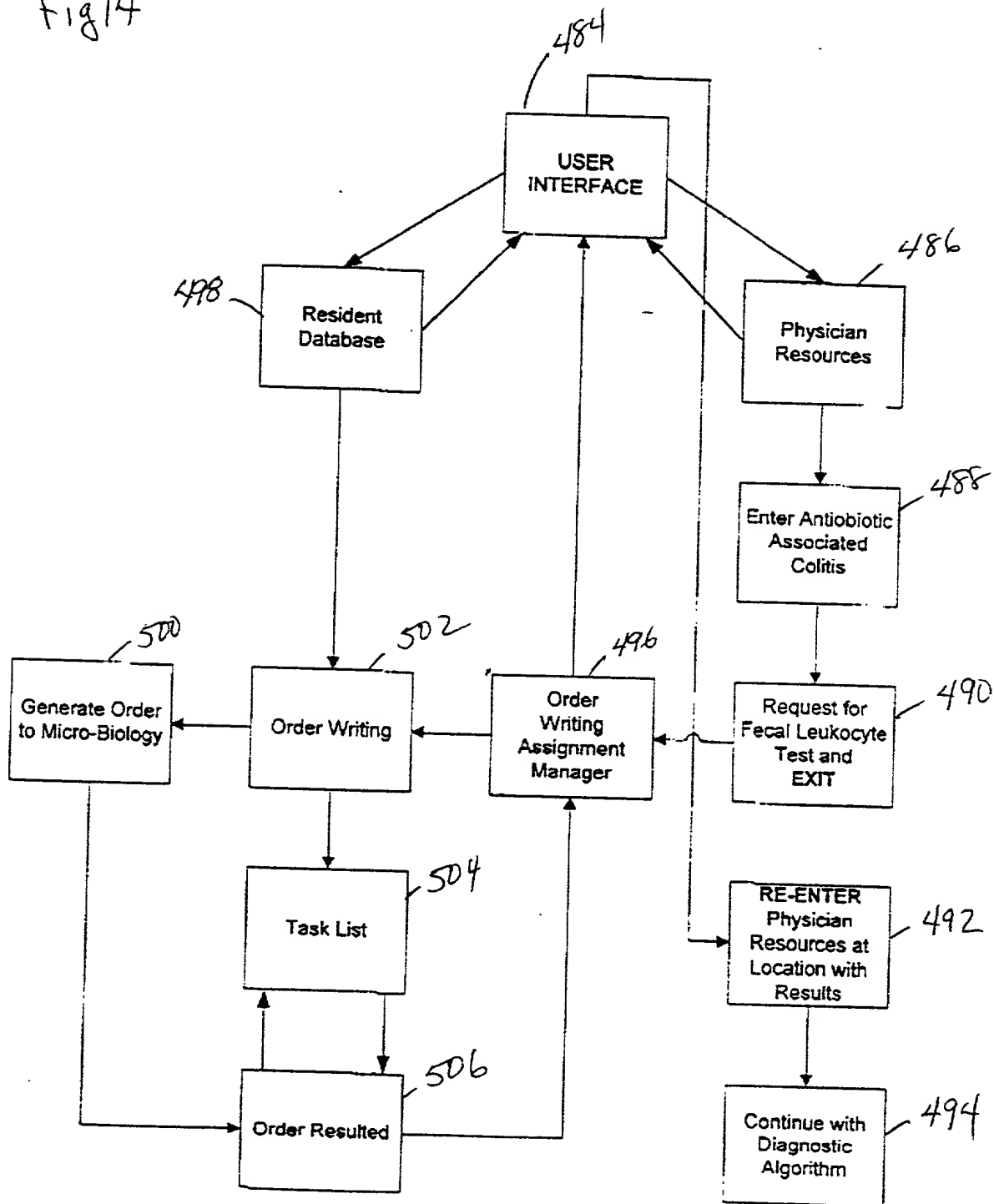


Fig 13B

09443072 111899

# PHYSICIAN RESOURCES AND ORDER WRITING DATA INTERFACE

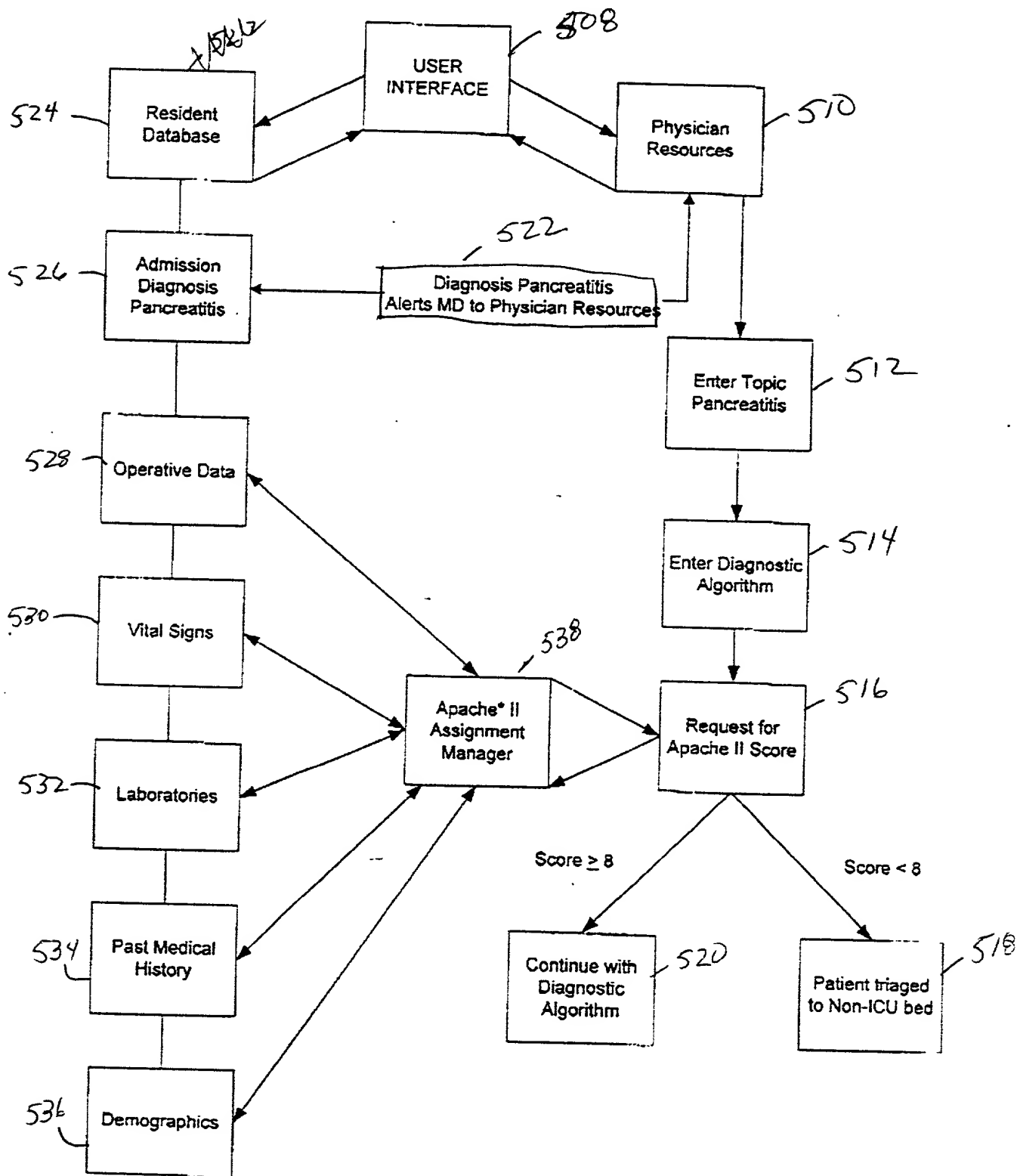
Fig 14



0944307 4489 65847 20E450

# PHYSICIAN RESOURCES DATABASE DATA INTERFACE

Fig 15



\*Assigns Apache II Score based upon weighted composite of 25 variables

6644307 24064450



# Order Writing Flow Sheet

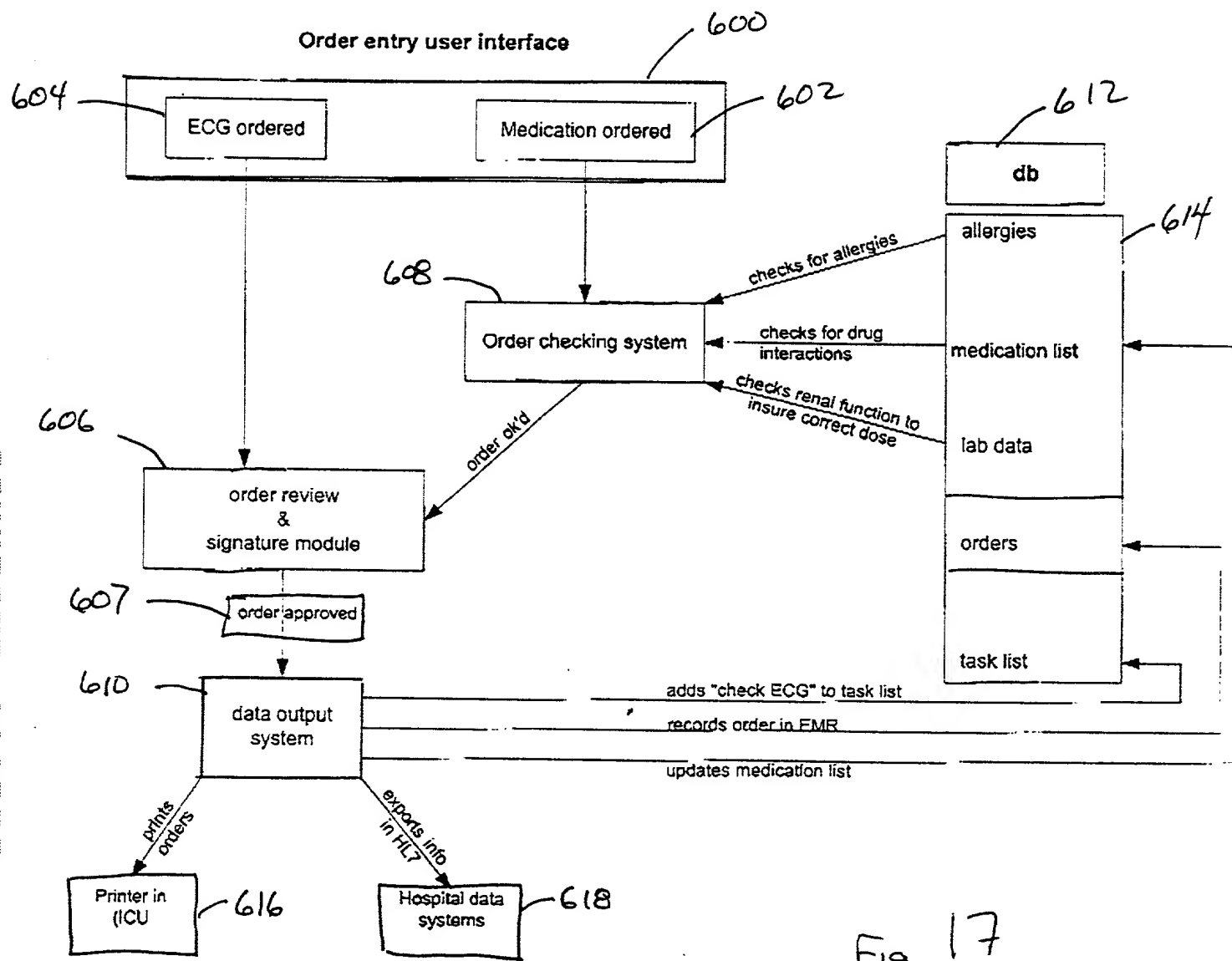


Fig 17

# Event Log

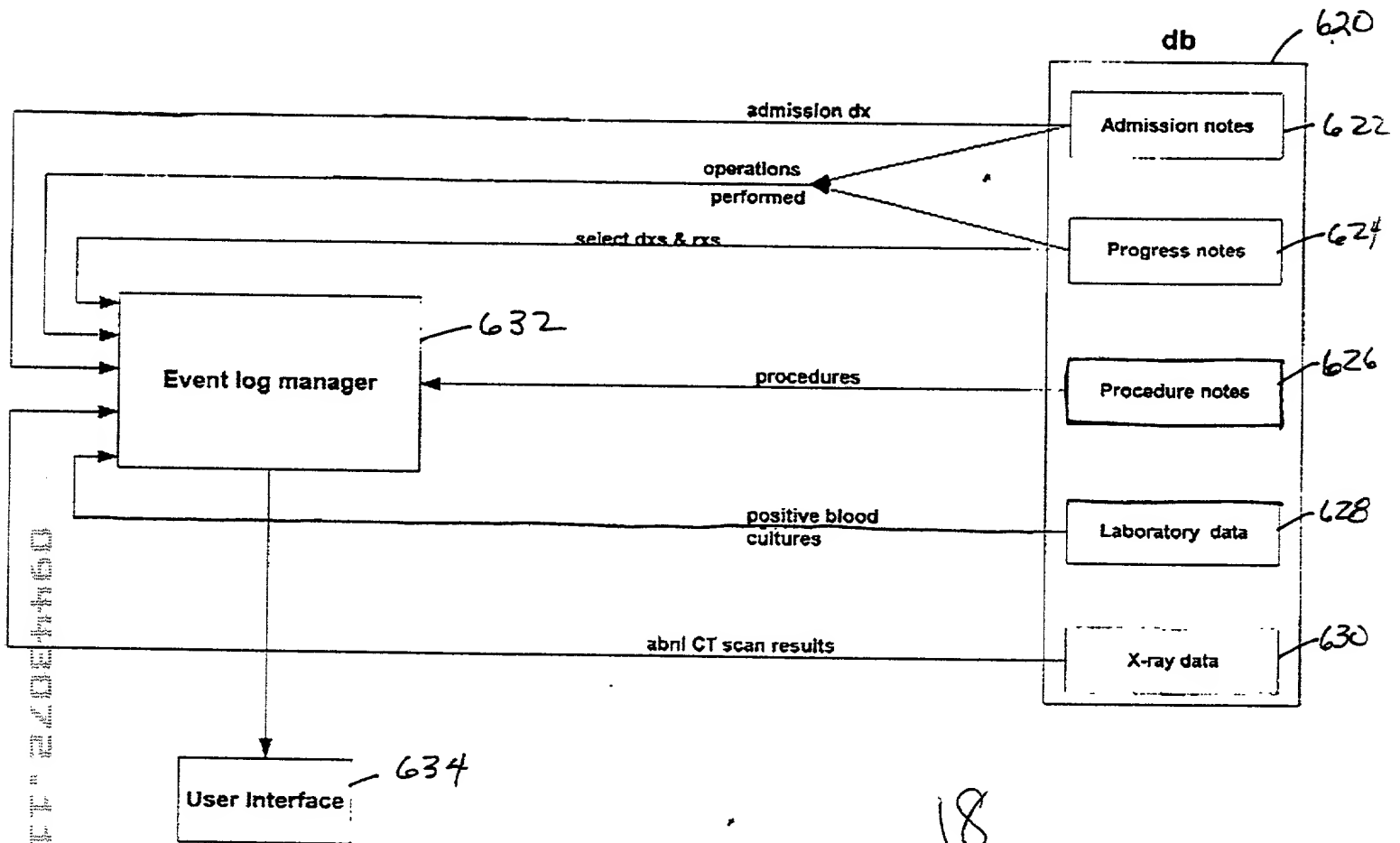
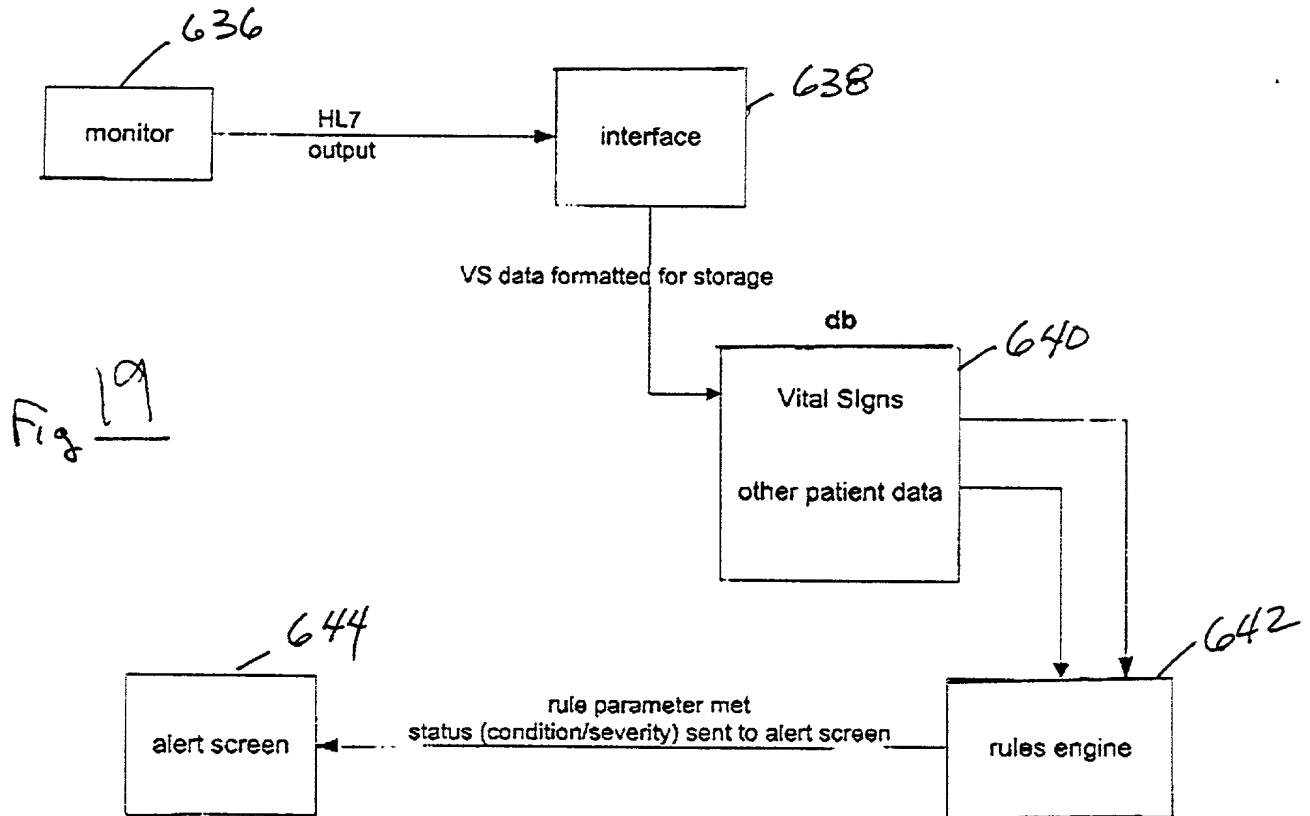


Fig 18

The event log presents in a single location key clinical information from throughout a patient's stay in the ICU. The event log provides care givers with a snapshot view of all salient events since admission. All relevant data are presented chronologically.

## SMART ALARMS

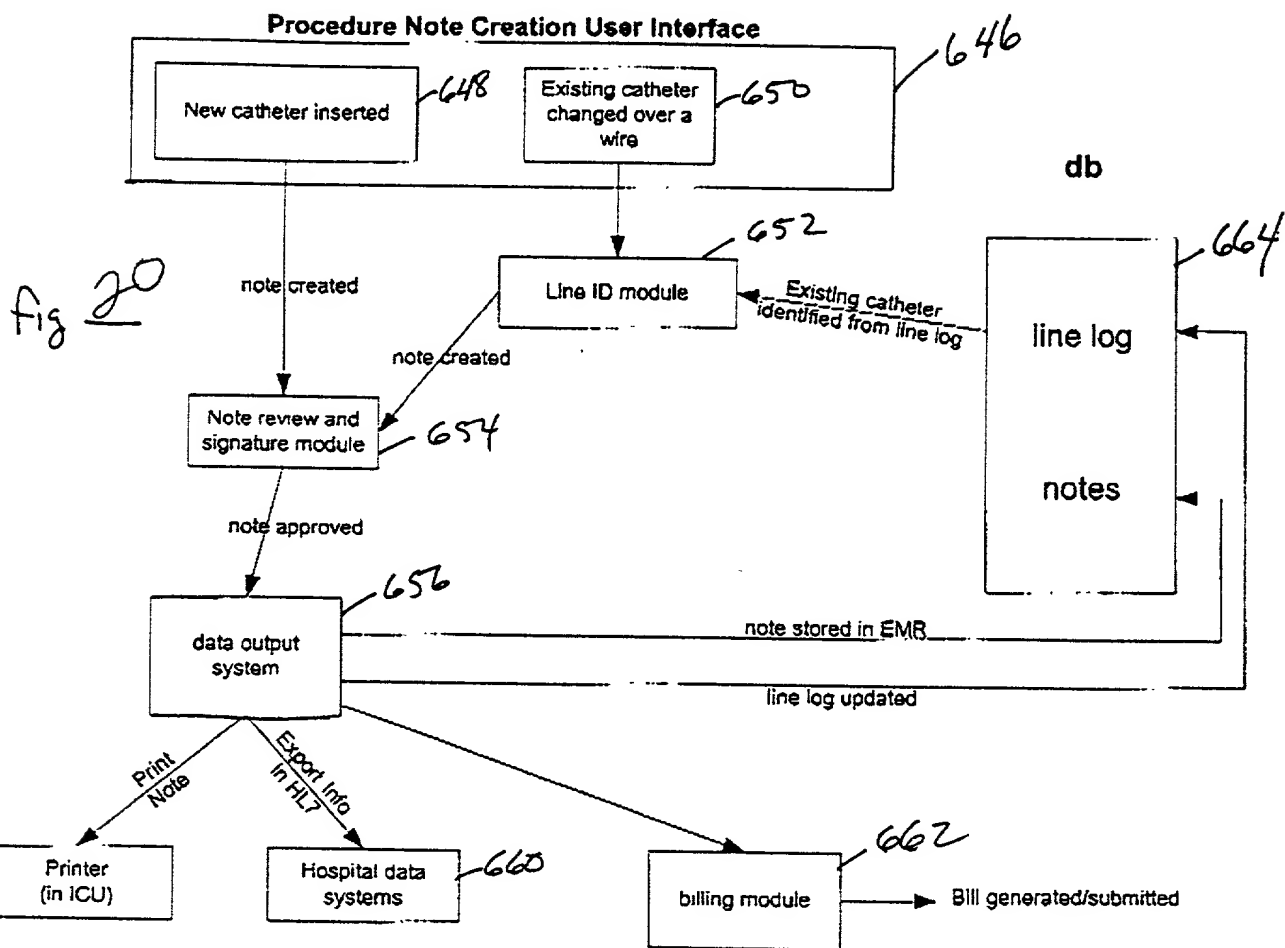


The smart alarm system constantly monitors physiologic data (collected once a minute from the bedside monitors) and other clinical information. The rules engine searches for patterns of data indicative of clinical deterioration. Examples include changes in vital signs over time (e.g. a 25% increase in the HR and a 20% decrease in BP), parallel reductions in urine output and central venous pressure that suggest developing hypovolemia, and progressive reductions in hemoglobin concentration over time that indicate a need to exclude active bleeding (and a possible need to administer blood). When rule conditions are met, relevant information is displayed on the system "alert screen".

The rationale underlying smart alarms is to facilitate detection of impending problems and to automate problem detection.

The system balances alarm sensitivity and specificity in order to maximize the benefit of the alarms to the intensivist.

# Procedure Note - Line Log



The line log contains, for each patient, relevant information about all indwelling catheters, including type and location of catheter, insertion date, the most recent date that the catheter was changed over a wire, and the date the catheter was removed. This information helps clinicians evaluate the likelihood that a given catheter is infected and guides management.



## Figure 21

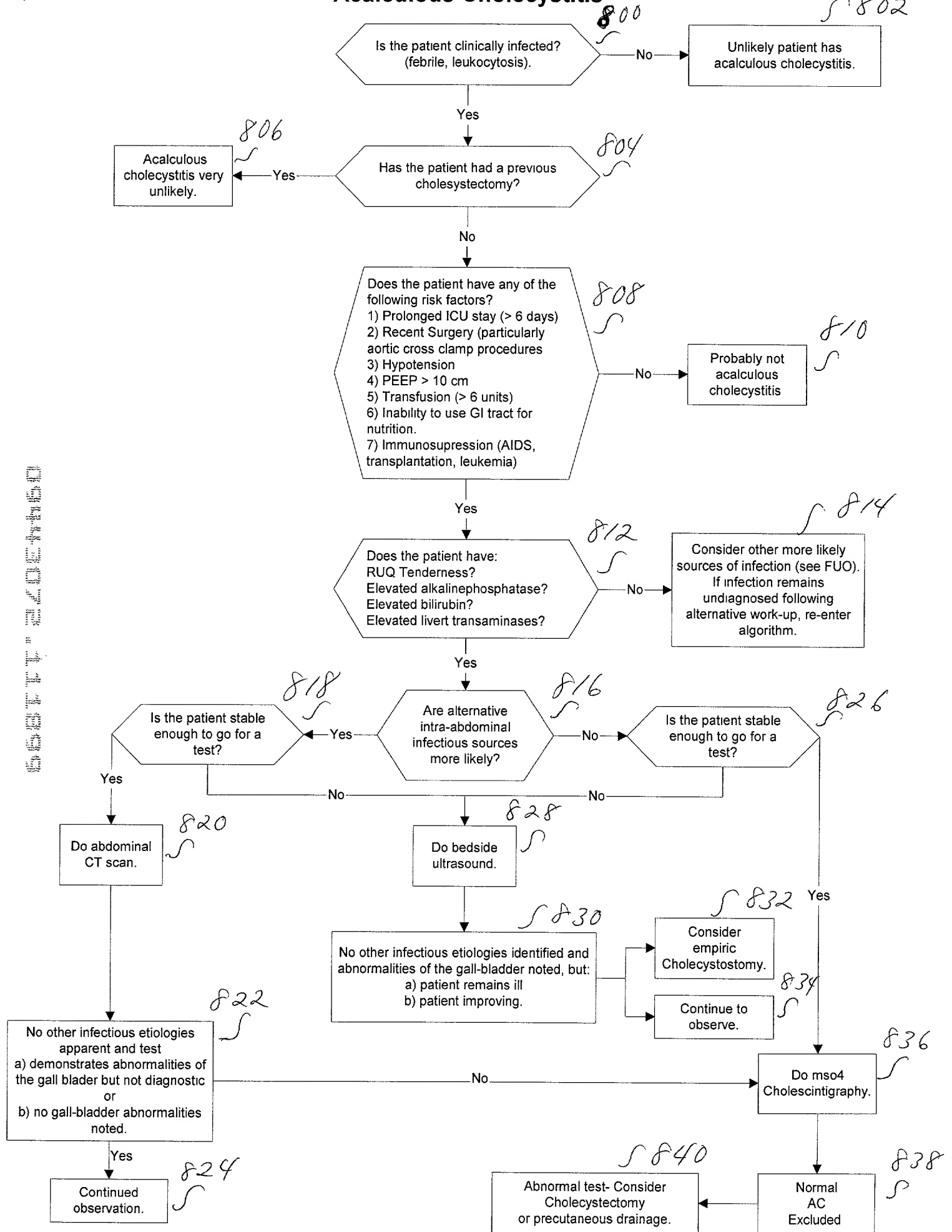
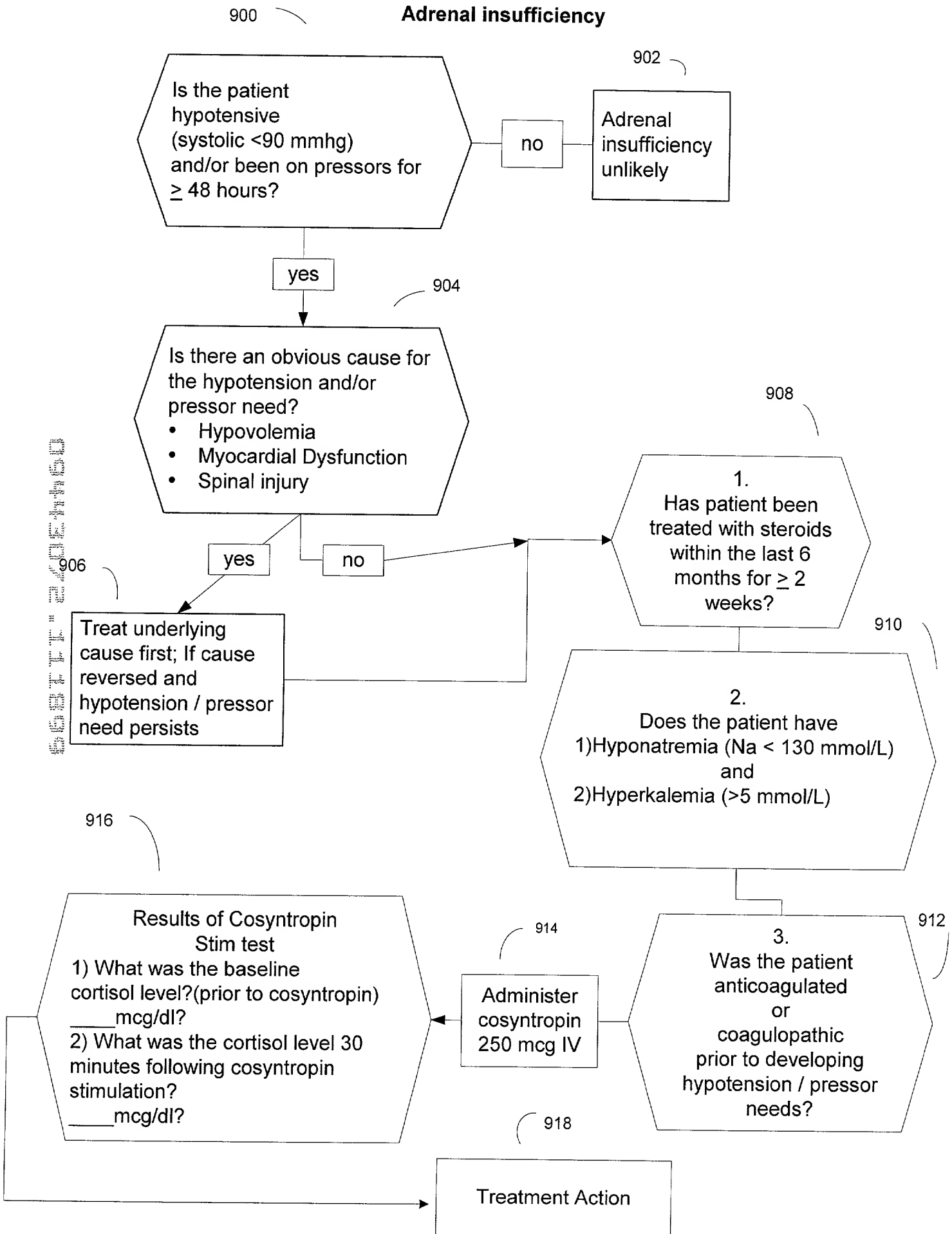
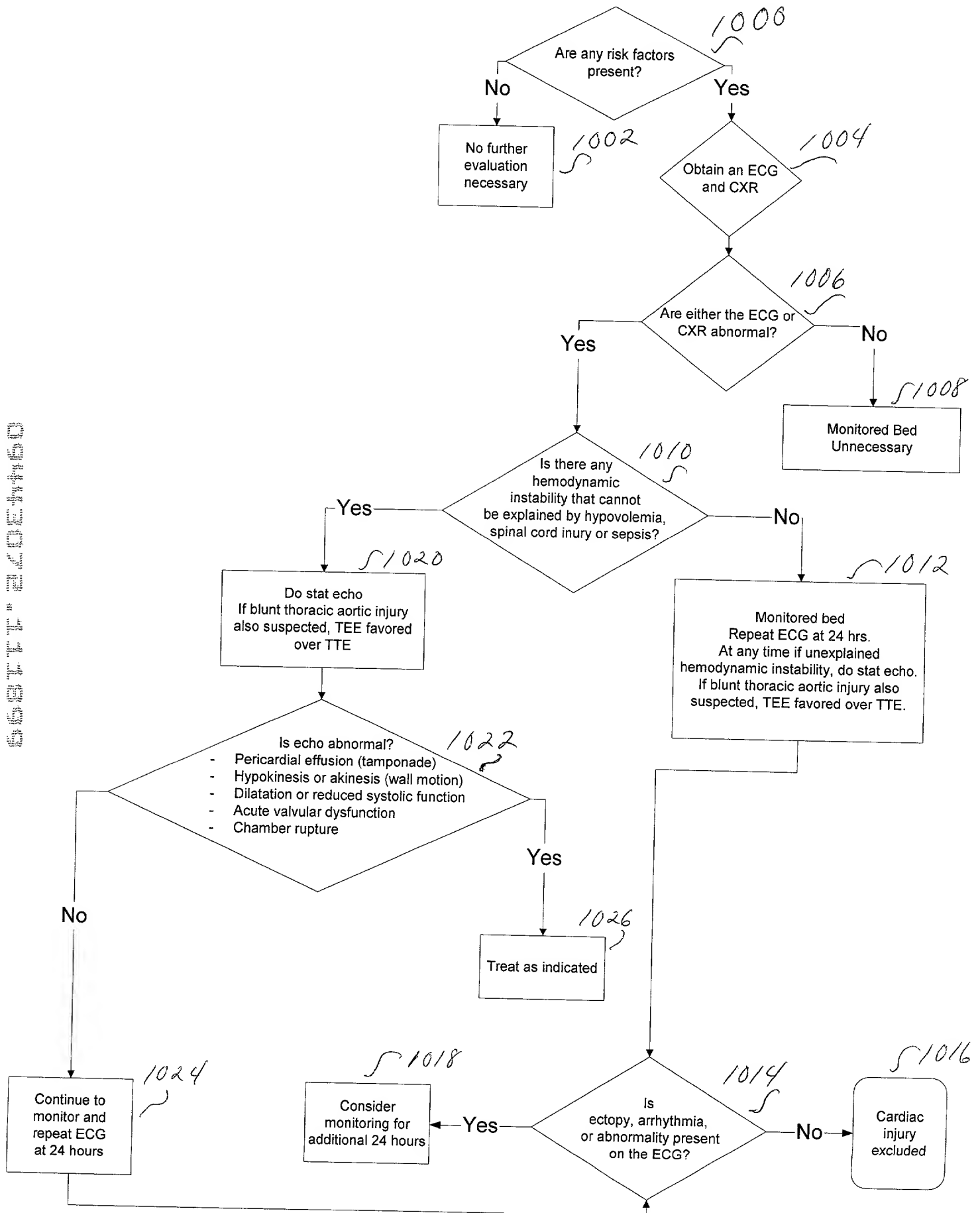


Figure 22

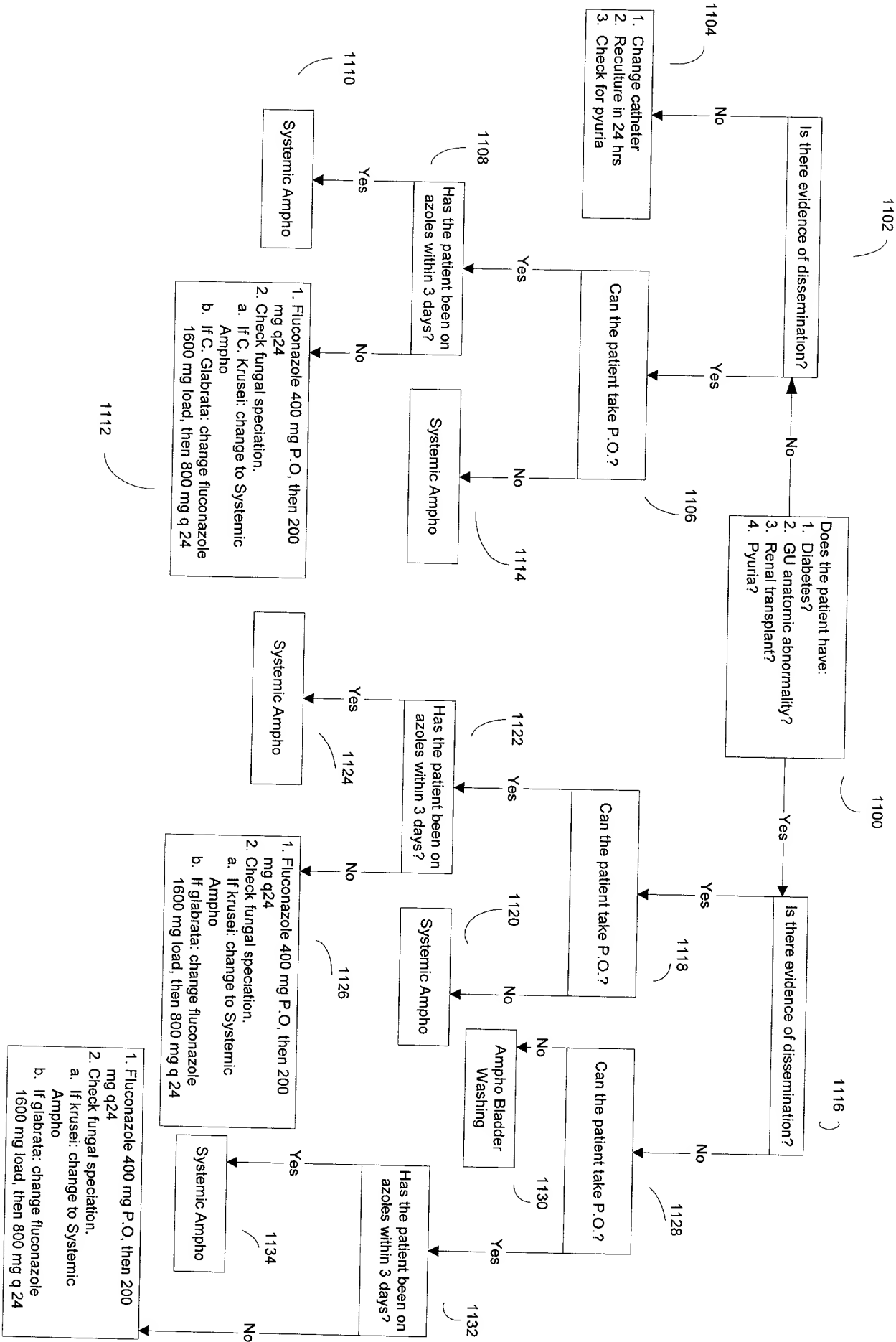


# Blunt Cardiac Injury

Figure 23

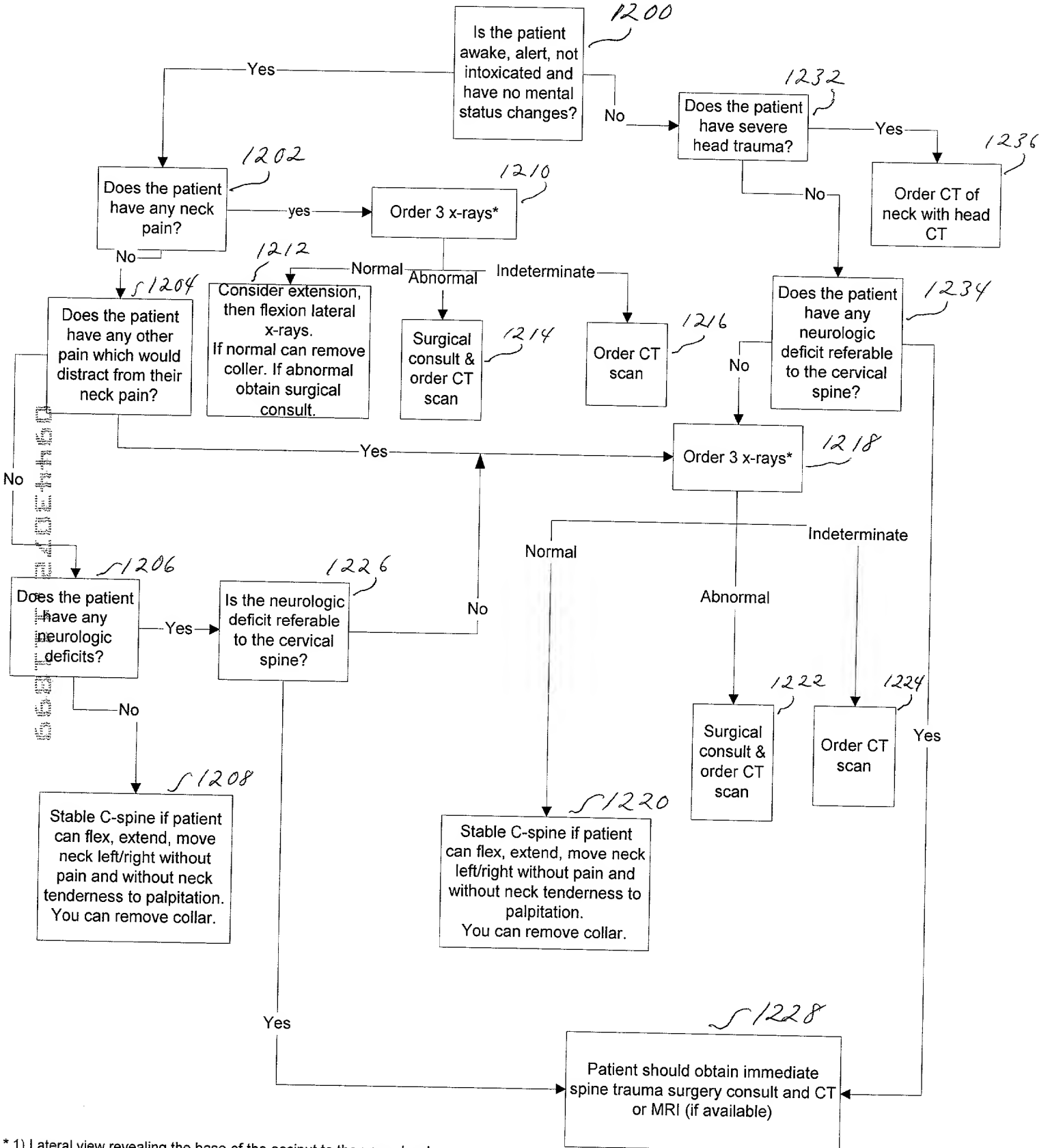


CANDIDURIA



0344 Figure 24-1-1399

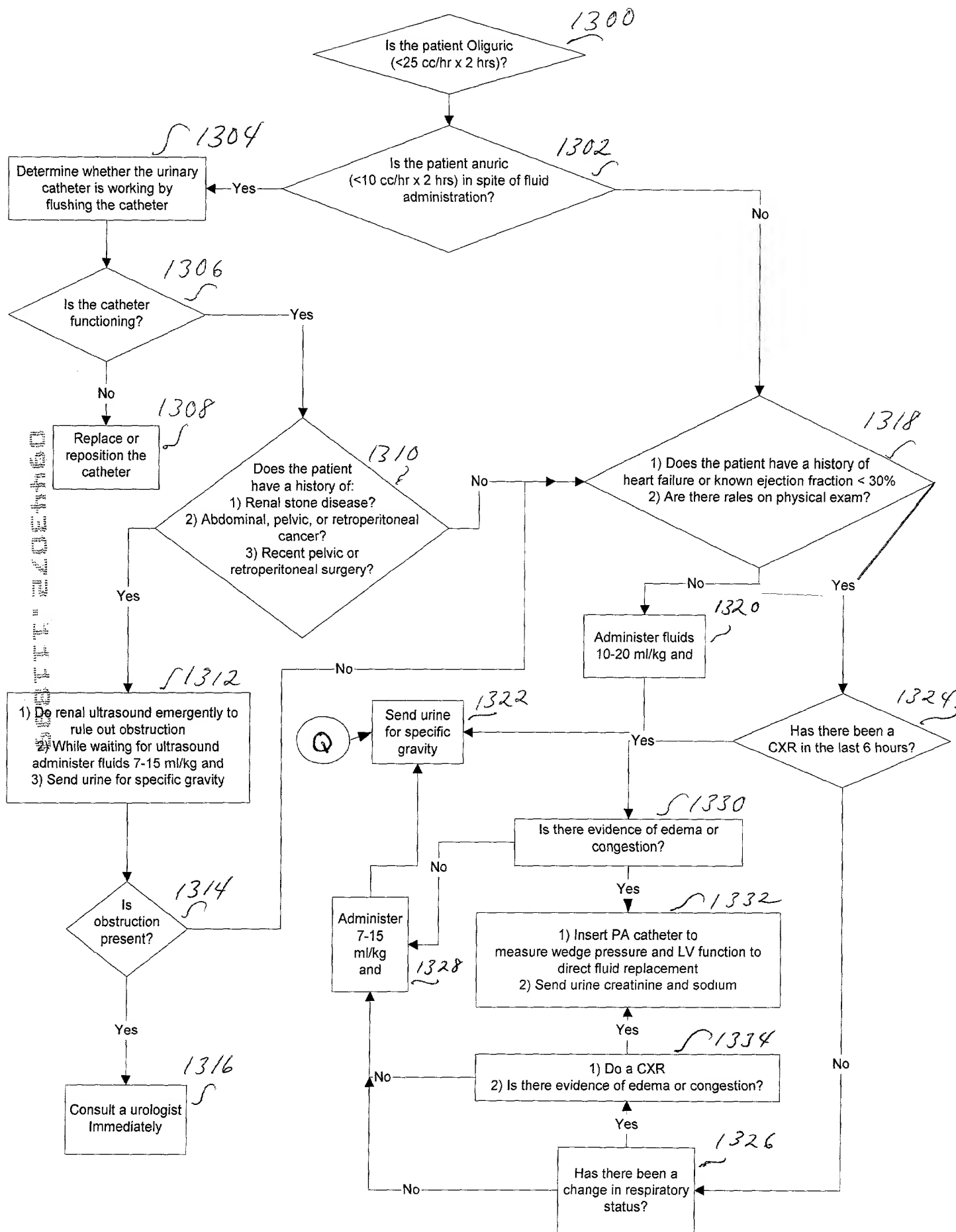
**Figure 25**



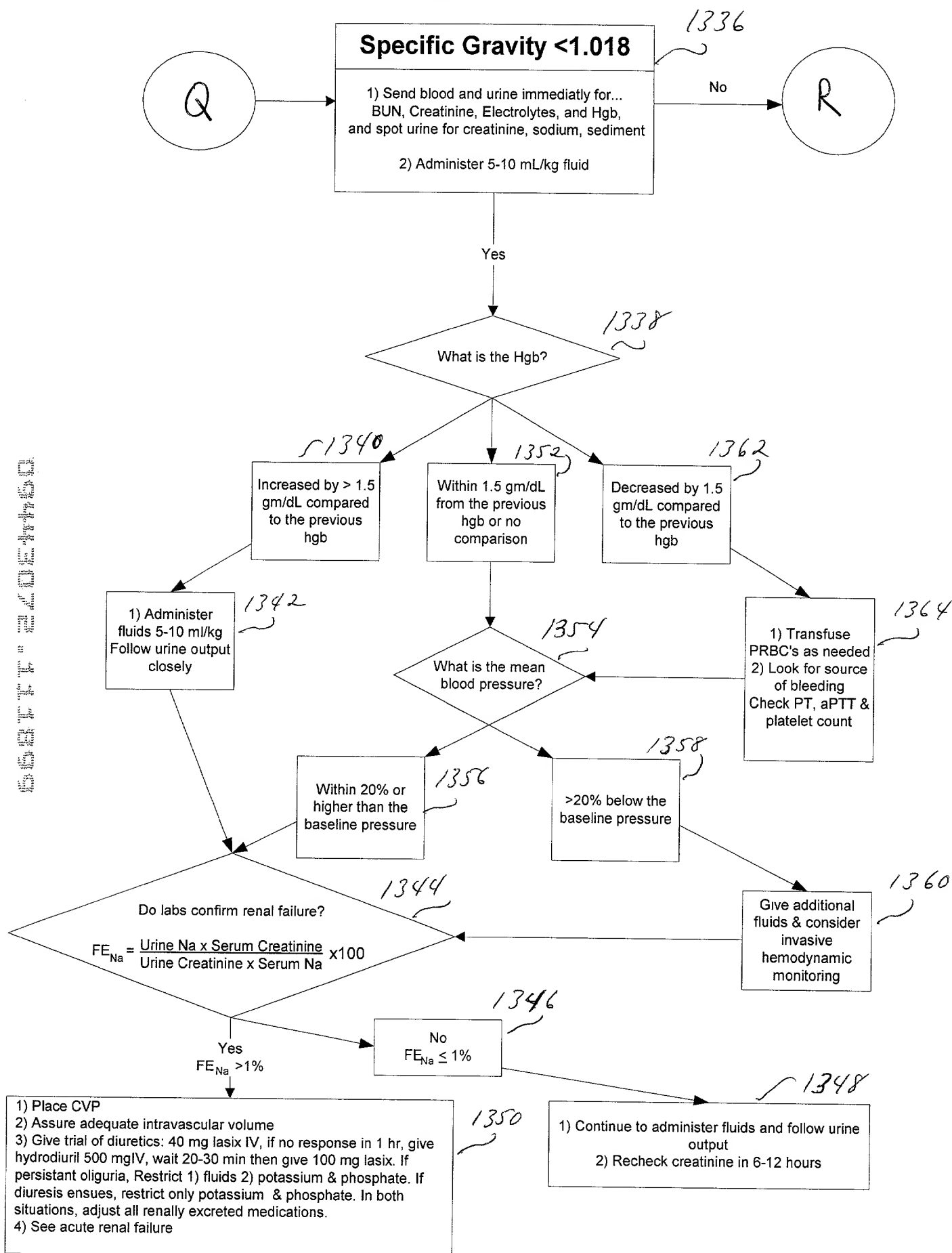
\* 1) Lateral view revealing the base of the occiput to the upper border of the first thoracic vertebra, 2) anteroposterior view revealing spinous processes of the second cervical through the first thoracic vertebra, and 3) an open mouth odontoid view revealing the lateral masses of the first cervical vertebra and entire odontoid process.

## Oliguria (page 1)

Figure 26



## Oliguria (page 2)



Oliguria (page 3)

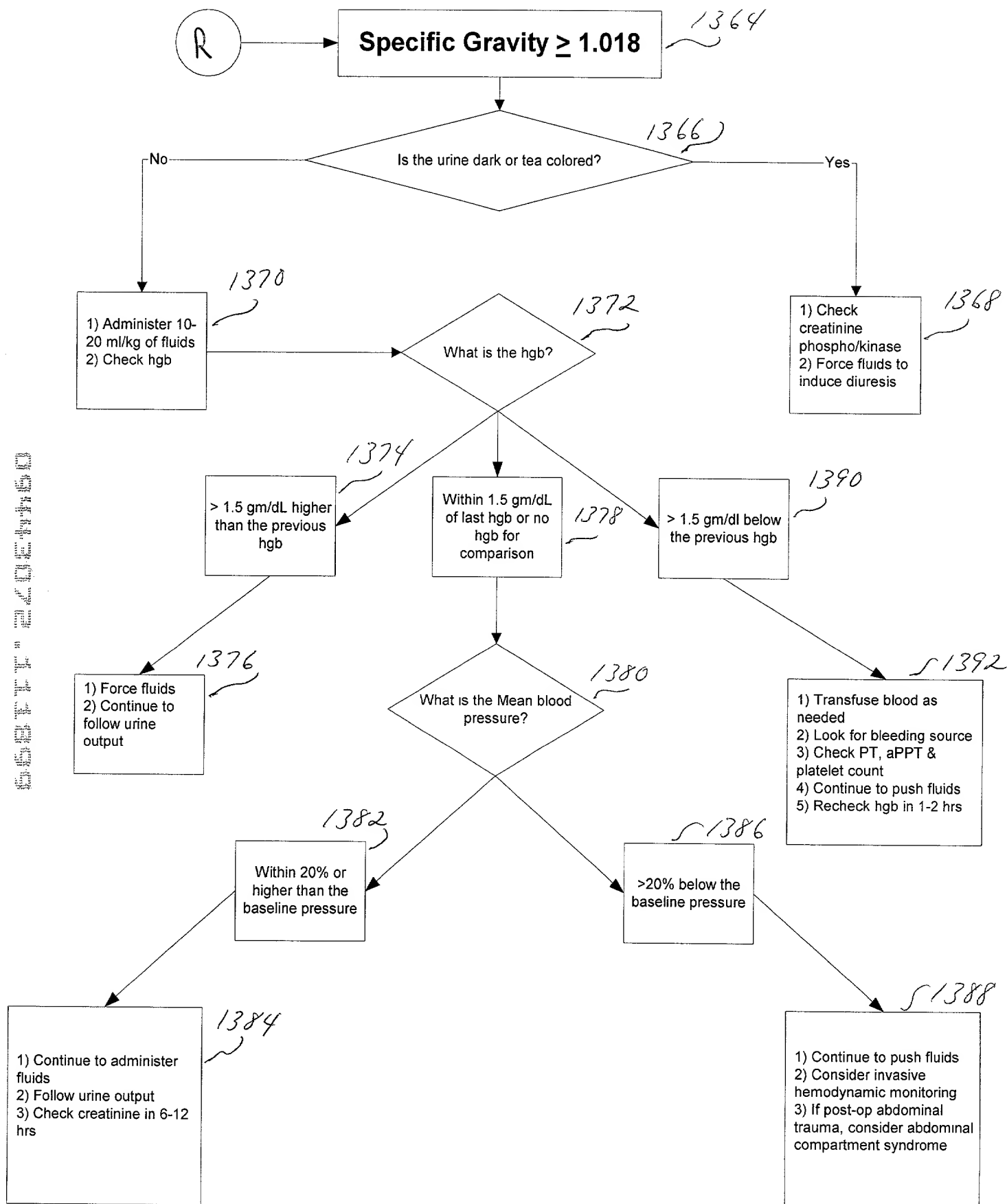
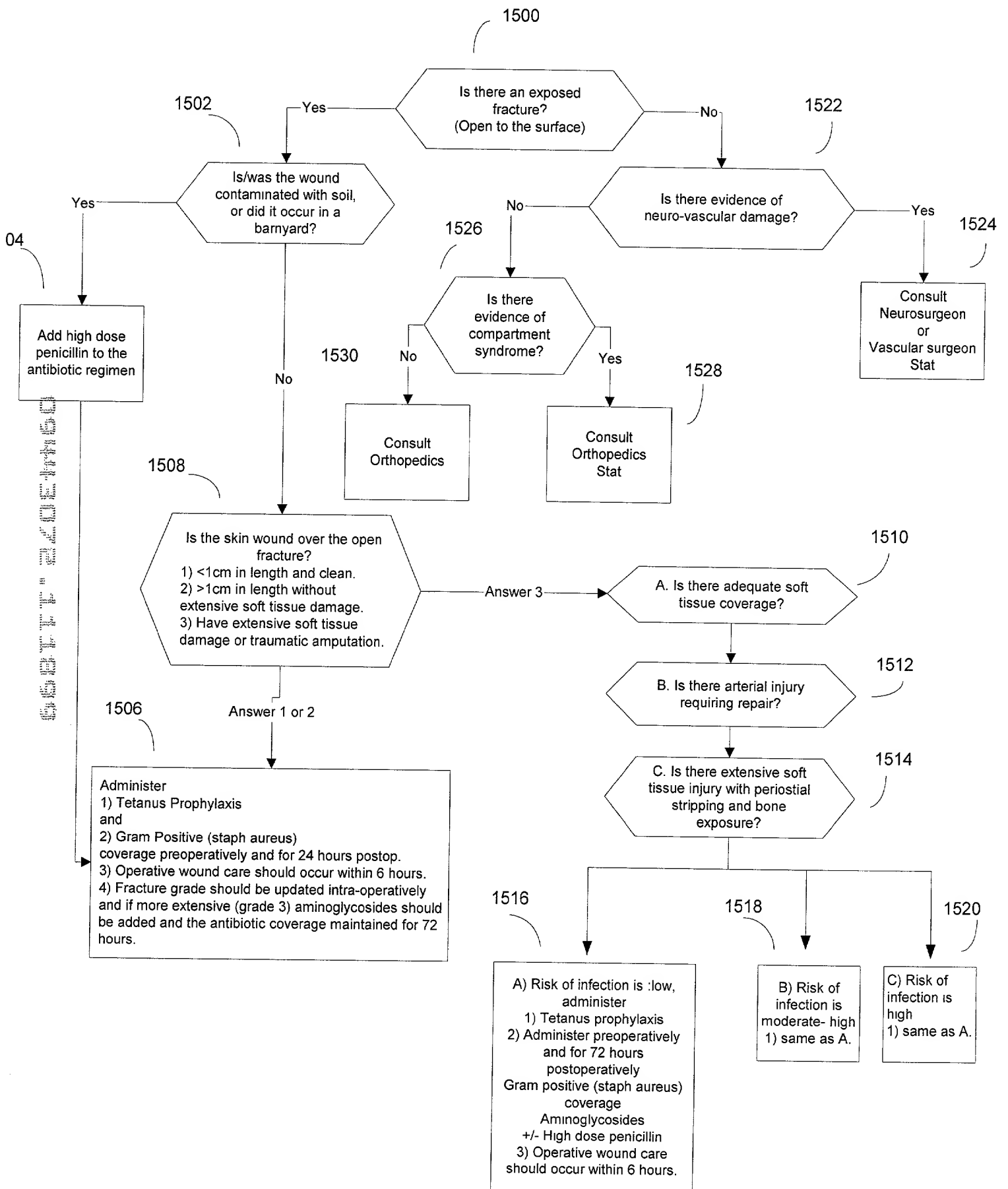




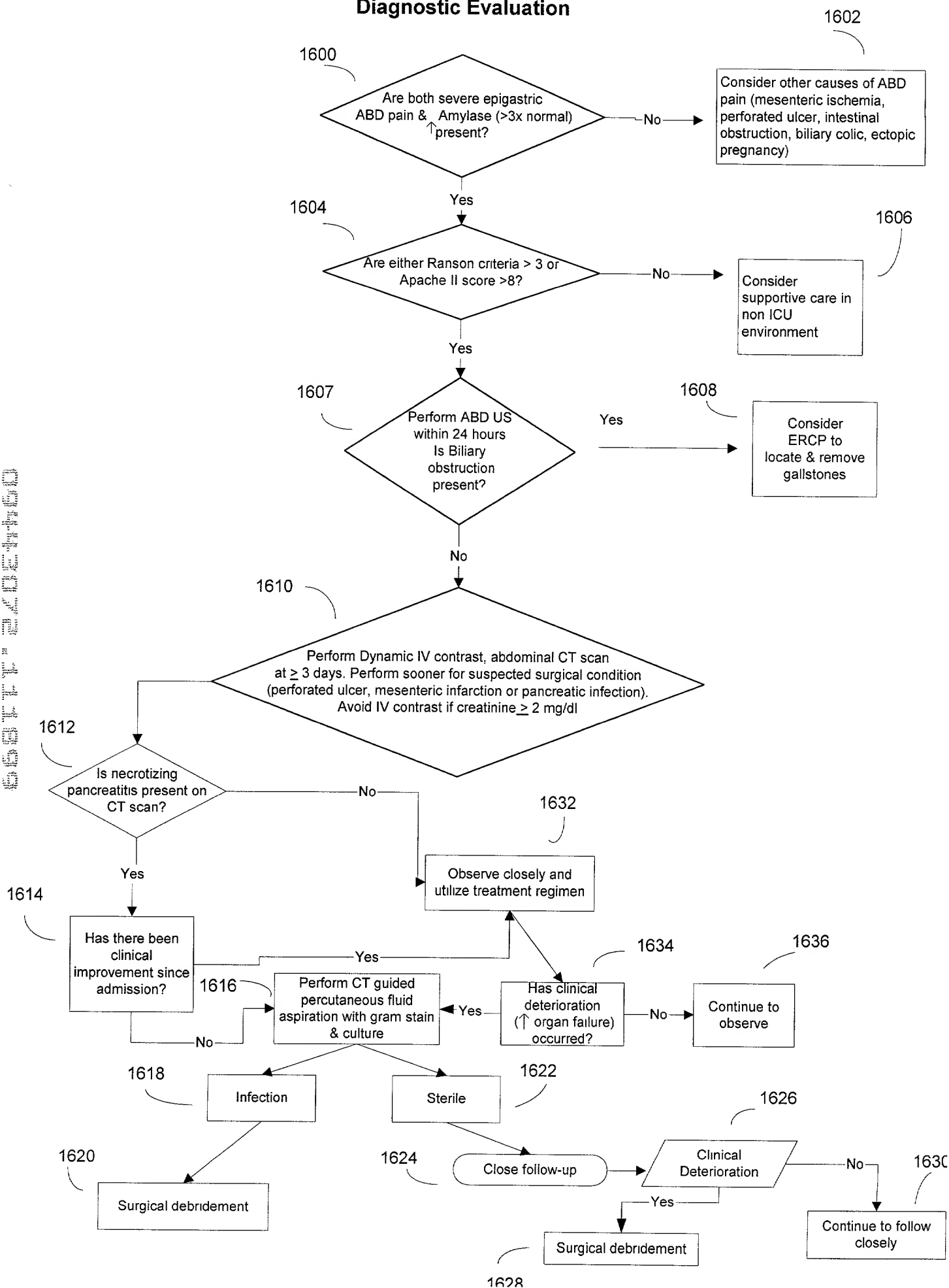
Figure 27

# OPEN FRACTURES



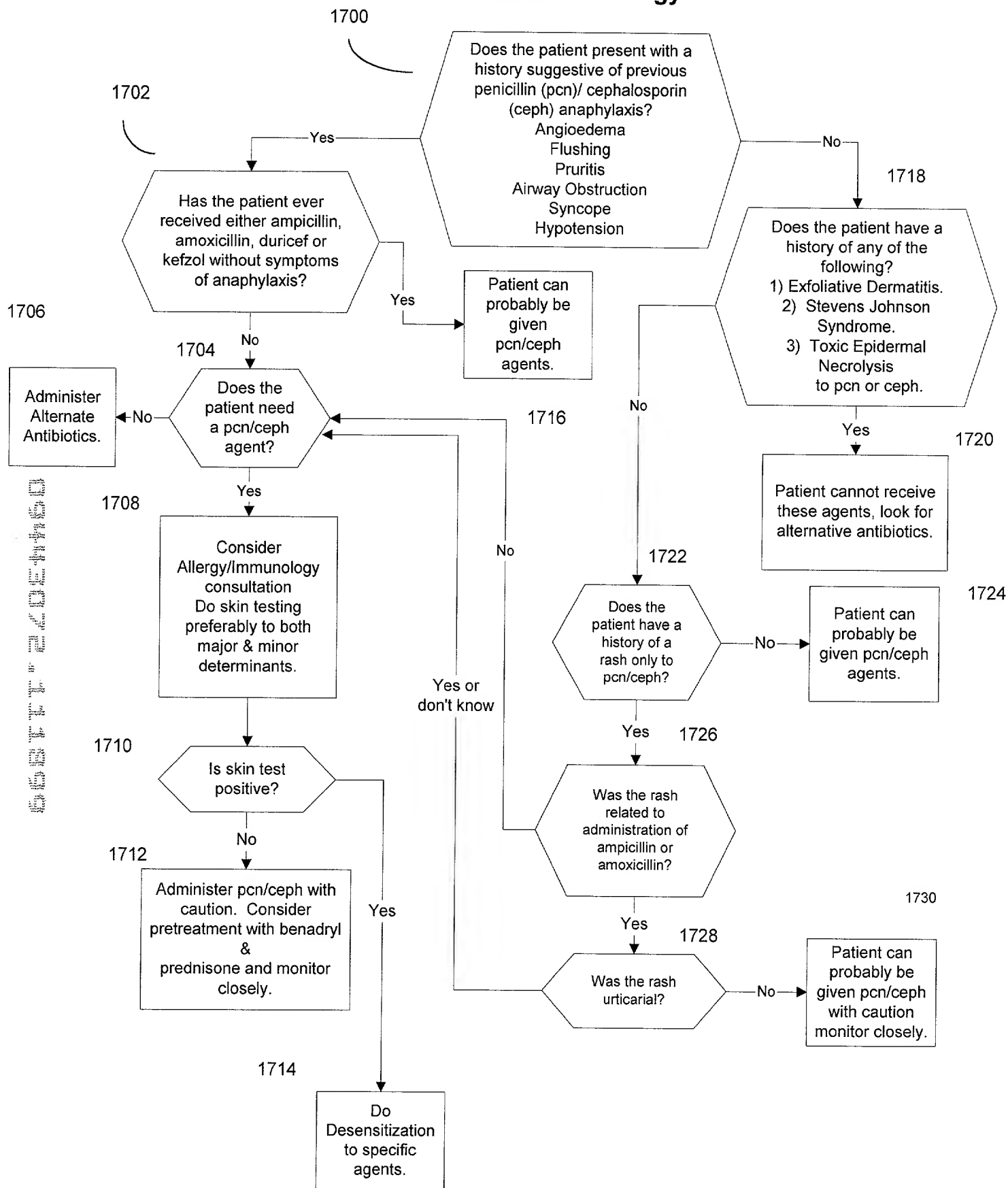
# **PANCREATITIS** **Diagnostic Evaluation**

65811" 24E4460



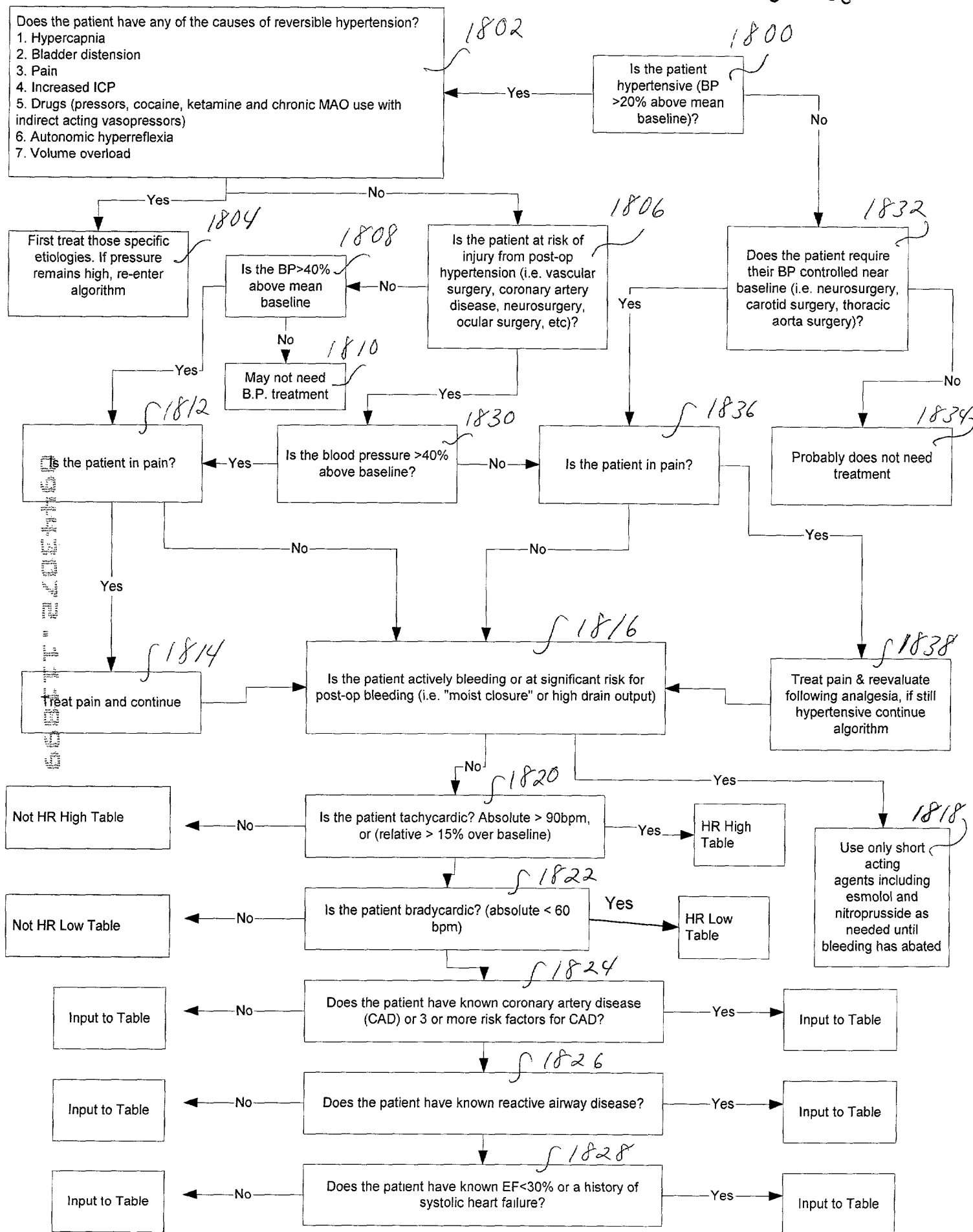
# Penicillin Allergy

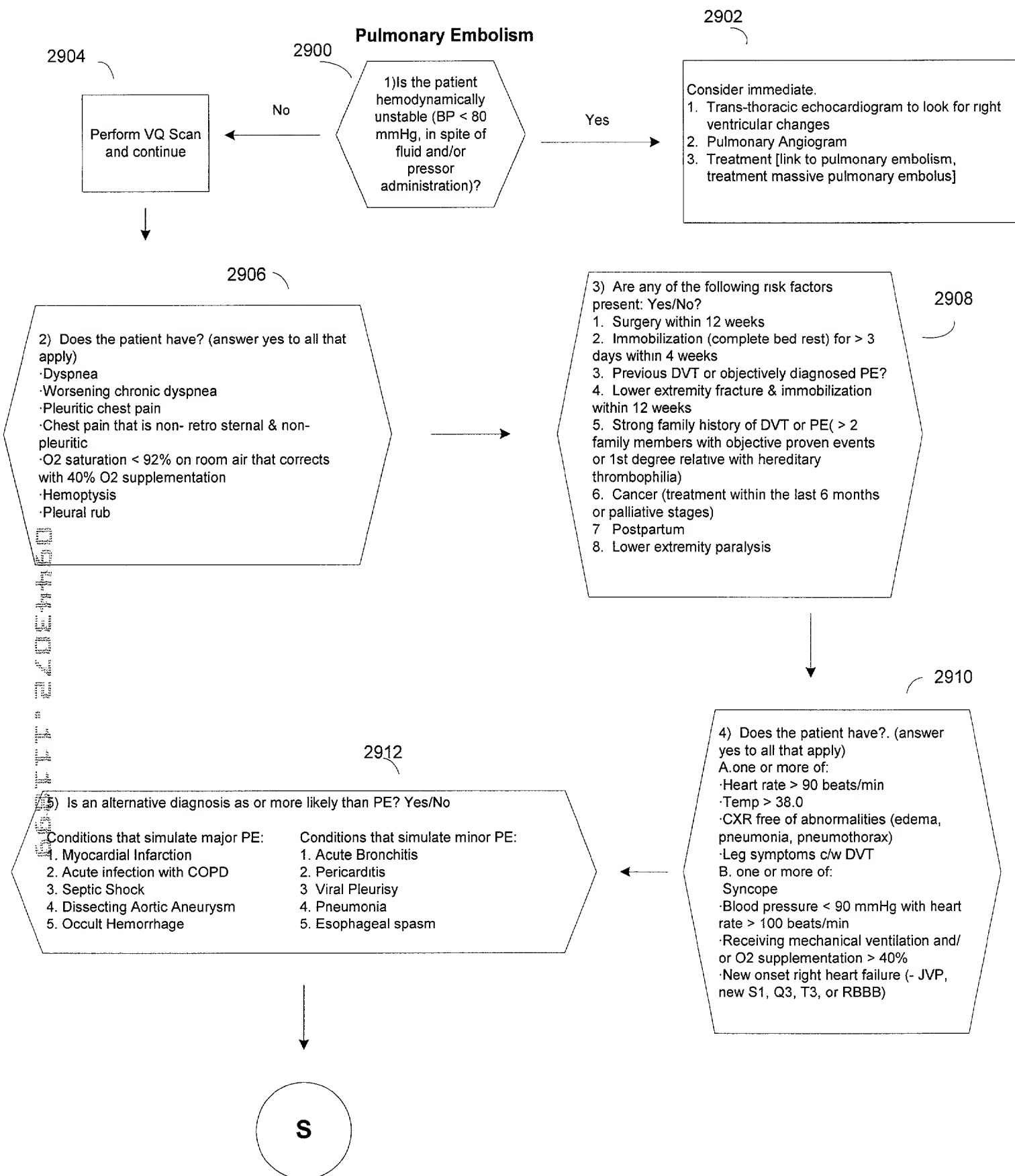
Figure 29



# Post-Op Hypertension

Figure 30





**Figure 31**

Pulmonary Embolism

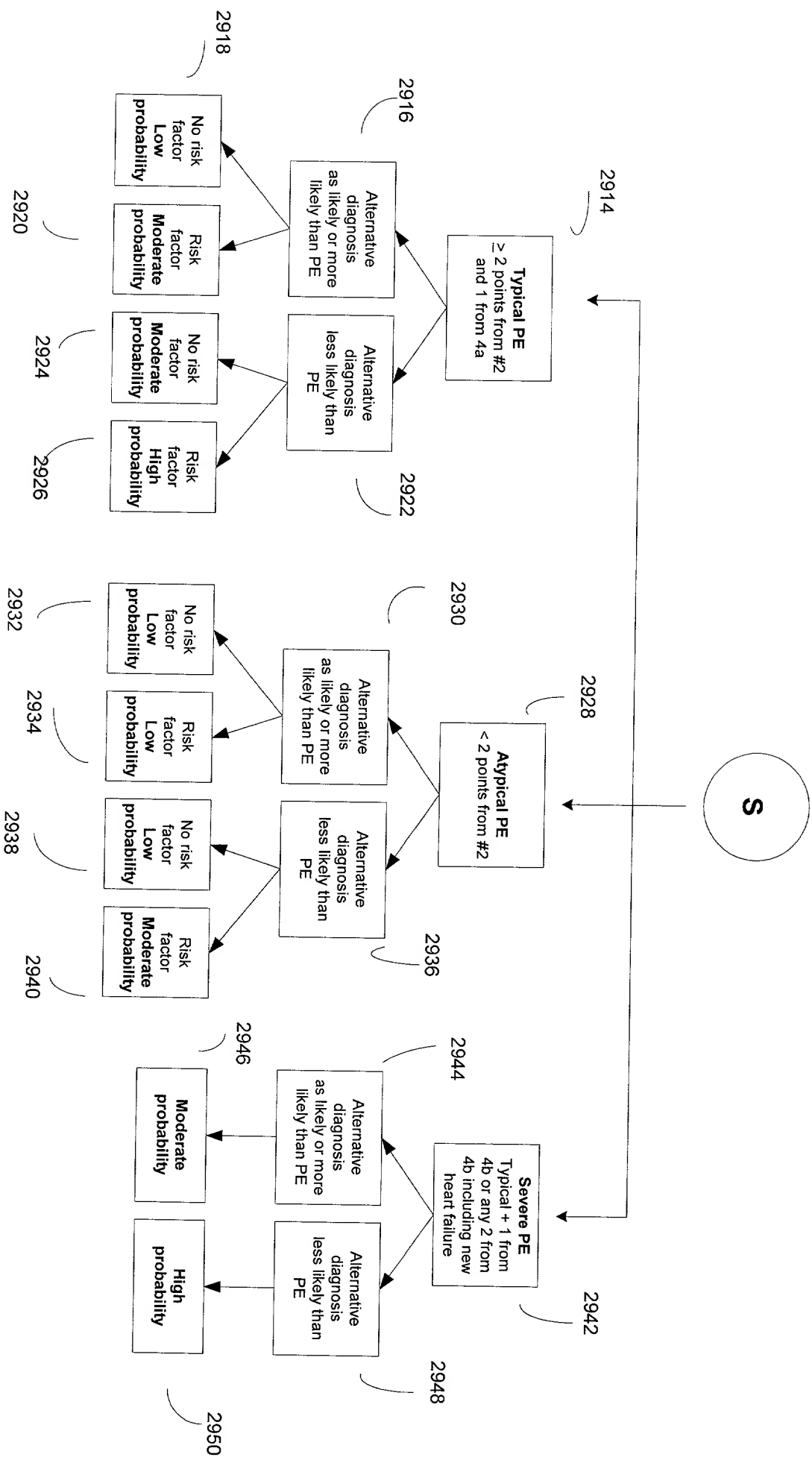


Figure 31A

0944303P 441893

# Seizure Algorithm

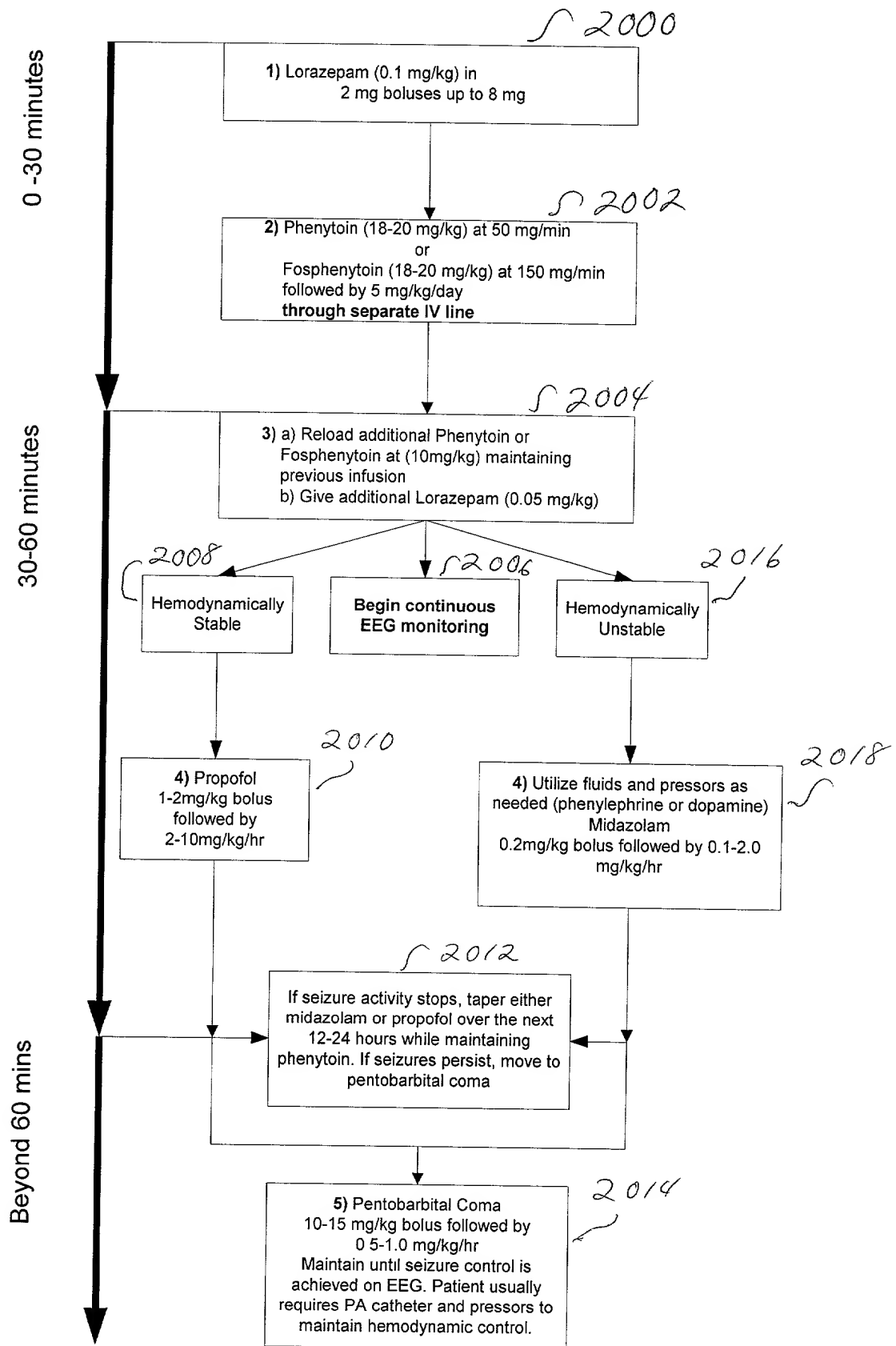


Figure 32

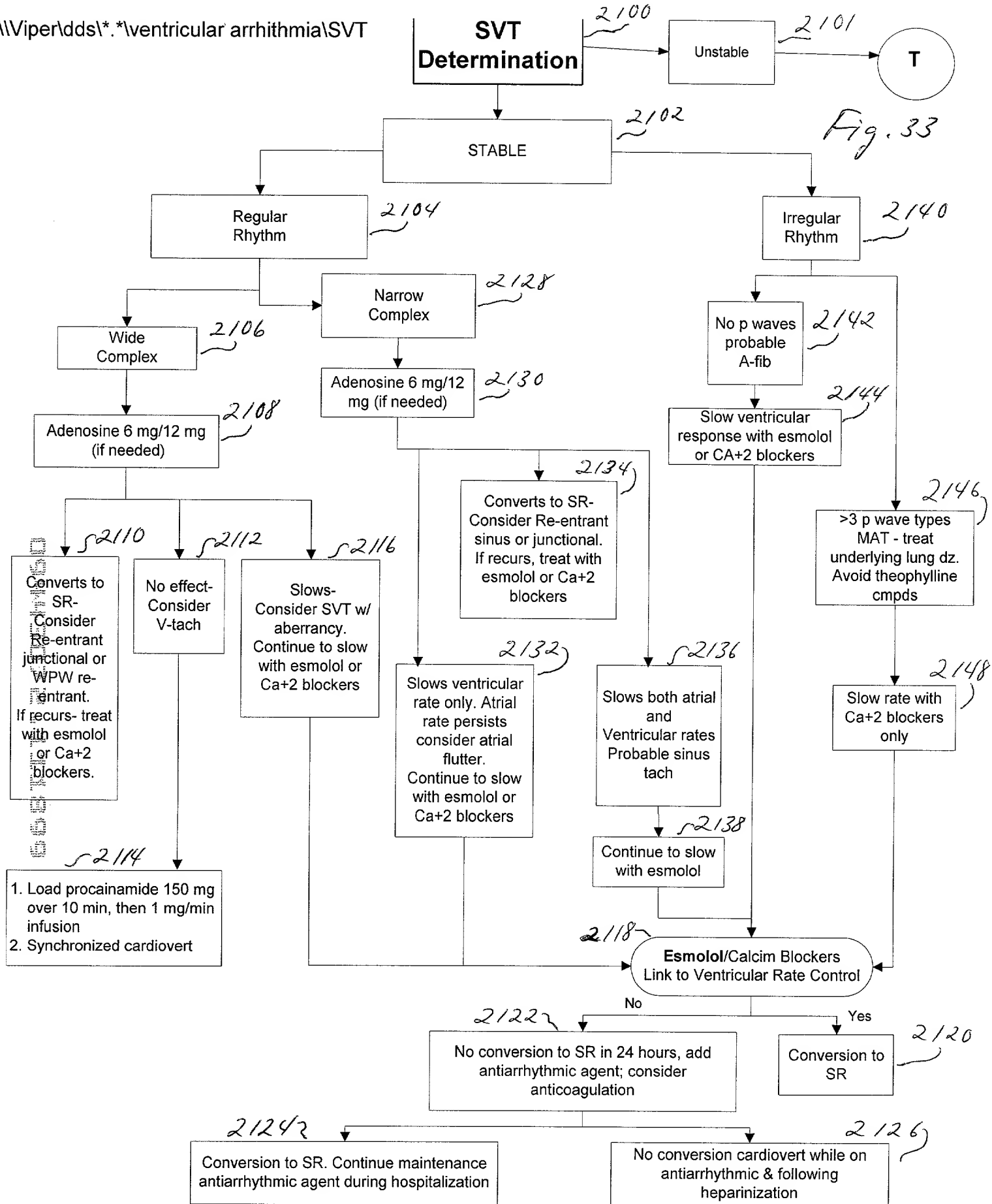
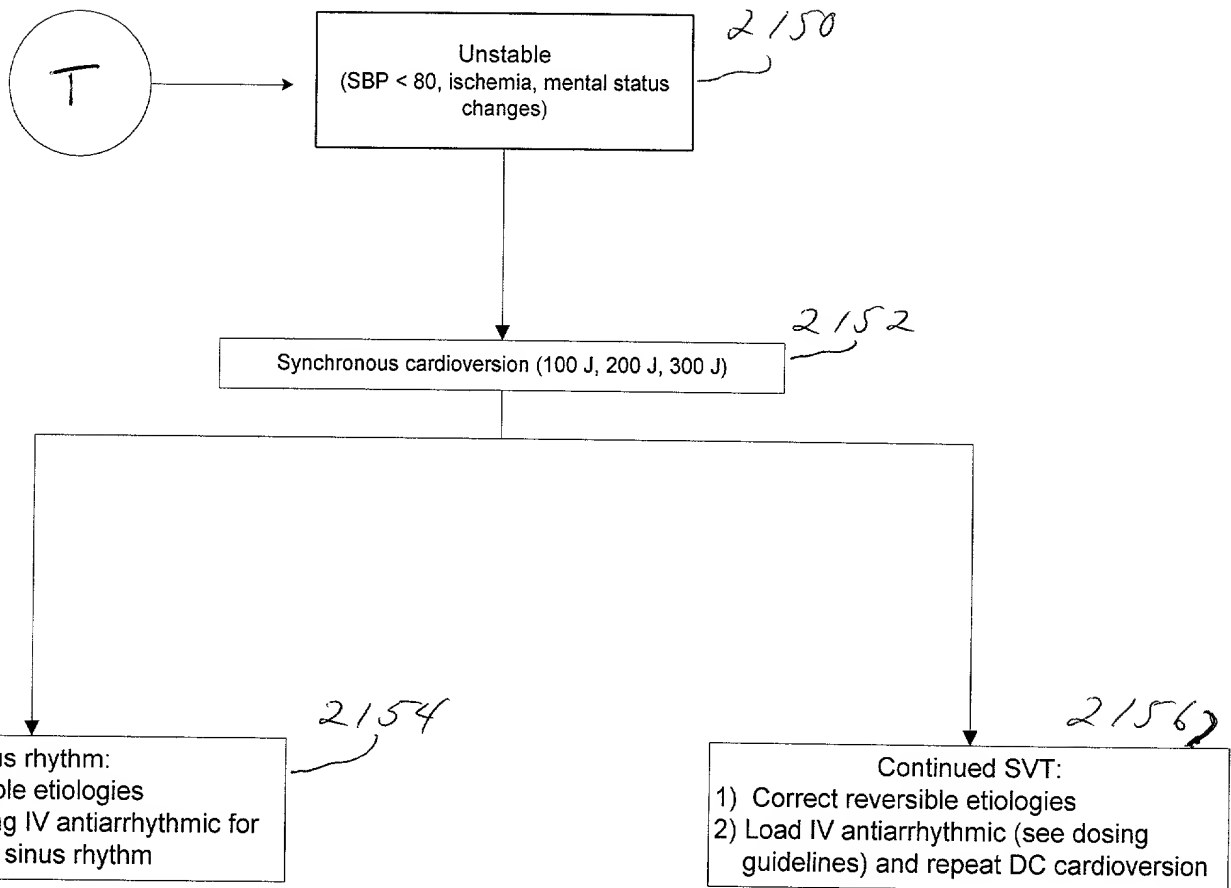


Figure 33



Fig 33A

# SVT Unstable



# Wide Complex QRS Tachycardia

Fig. 34

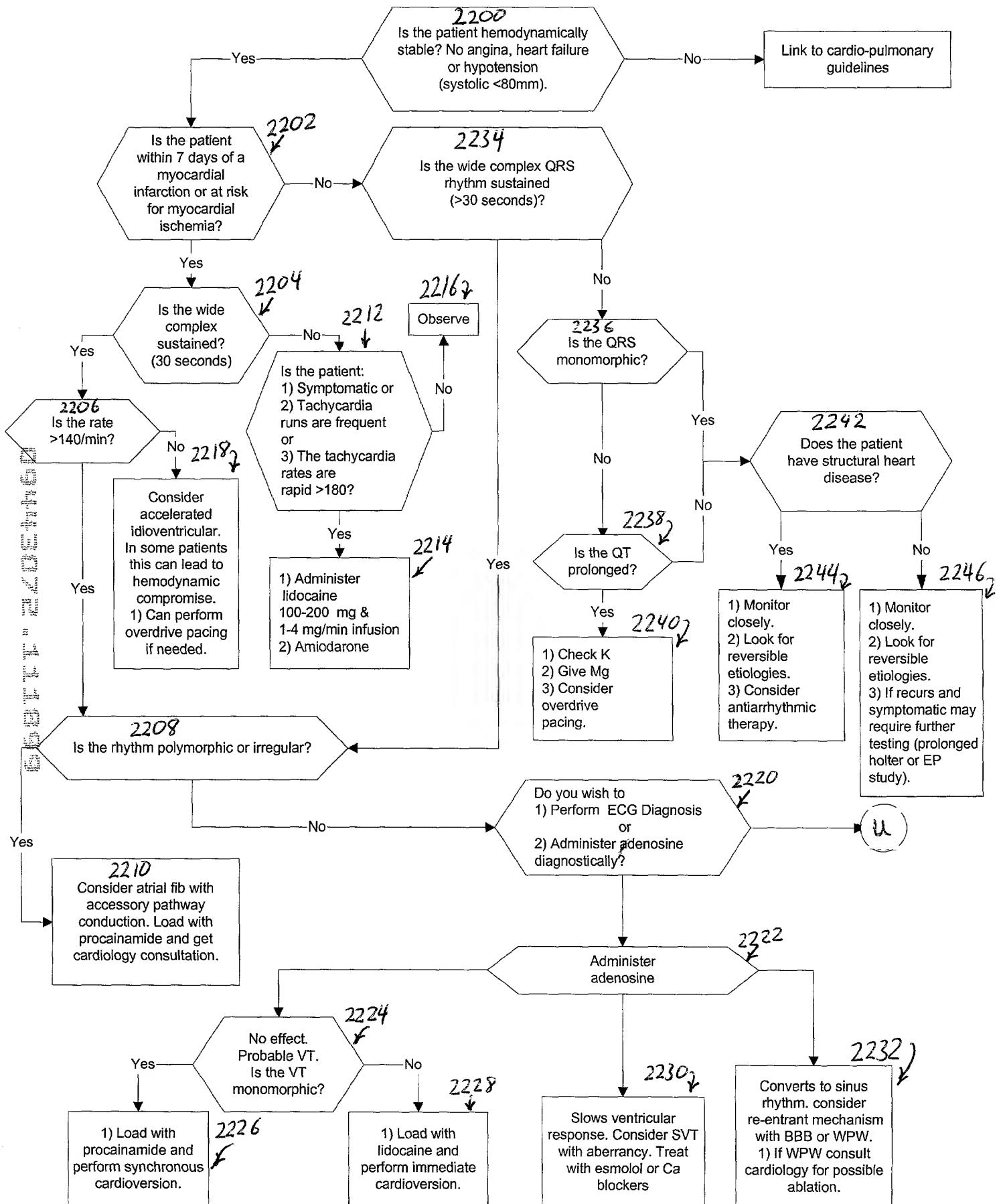


Fig. 34A

**Wide Complex QRS Tachycardia  
(page 2)  
ECG Diagnosis**

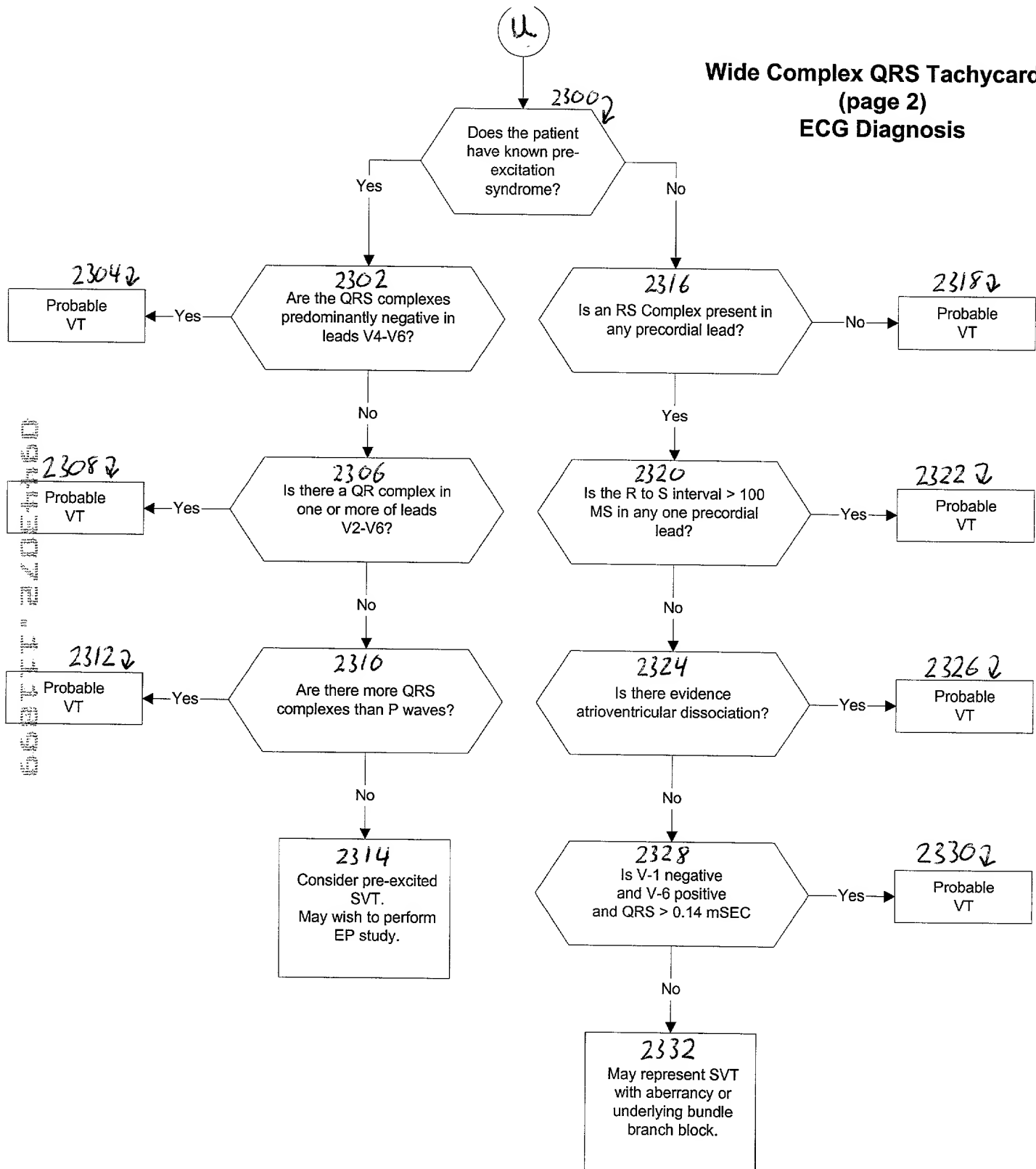
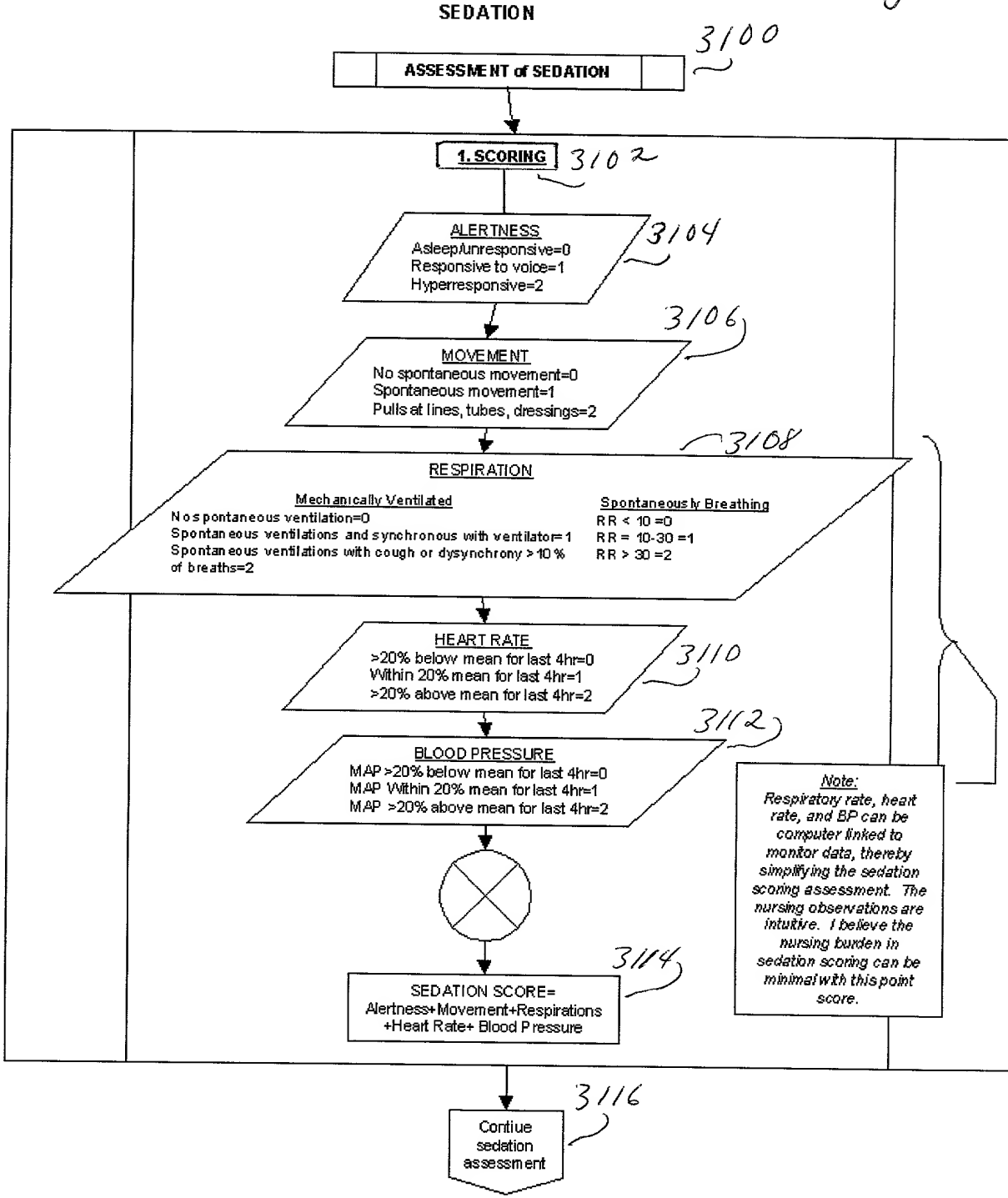
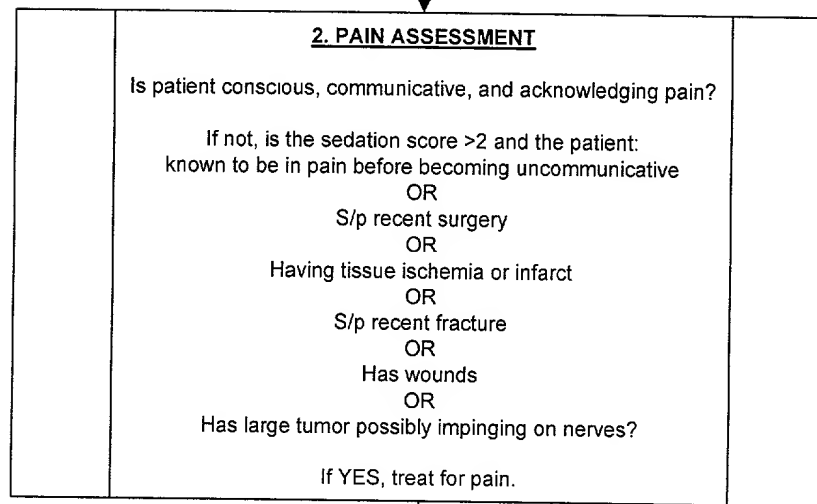


Fig. 41

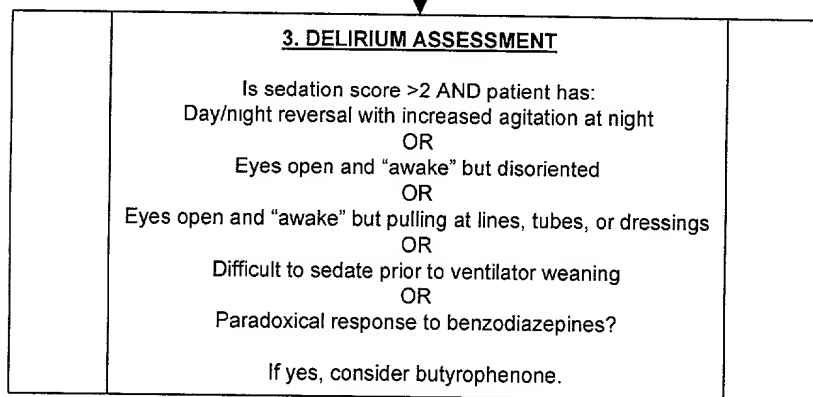


Sedation  
Assessment  
Continued

Figure 41A



3118



3120

Fig. 42

3200 ✓  
Bolus sliding scale  
midazolam



3202 ✓

If lorazepam <0-2 mg IV q 6hr then give midazolam 1-2 mg q 5min until adequately sedated  
If lorazepam =2-4 mg IV q 4hr then give midazolam 2 mg q 5min until adequately sedated  
If lorazepam =5-10 mg IV q 4hr then give midazolam 2-5 mg q 5min until adequately sedated  
If lorazepam >10 mg IV q 4hr then give midazolam 5 mg q 5min until adequately sedated AND  
consider fentanyl and/or droperidol or Haldol for synergy despite delirium and pain assessment

030443072 441899

Figure 43

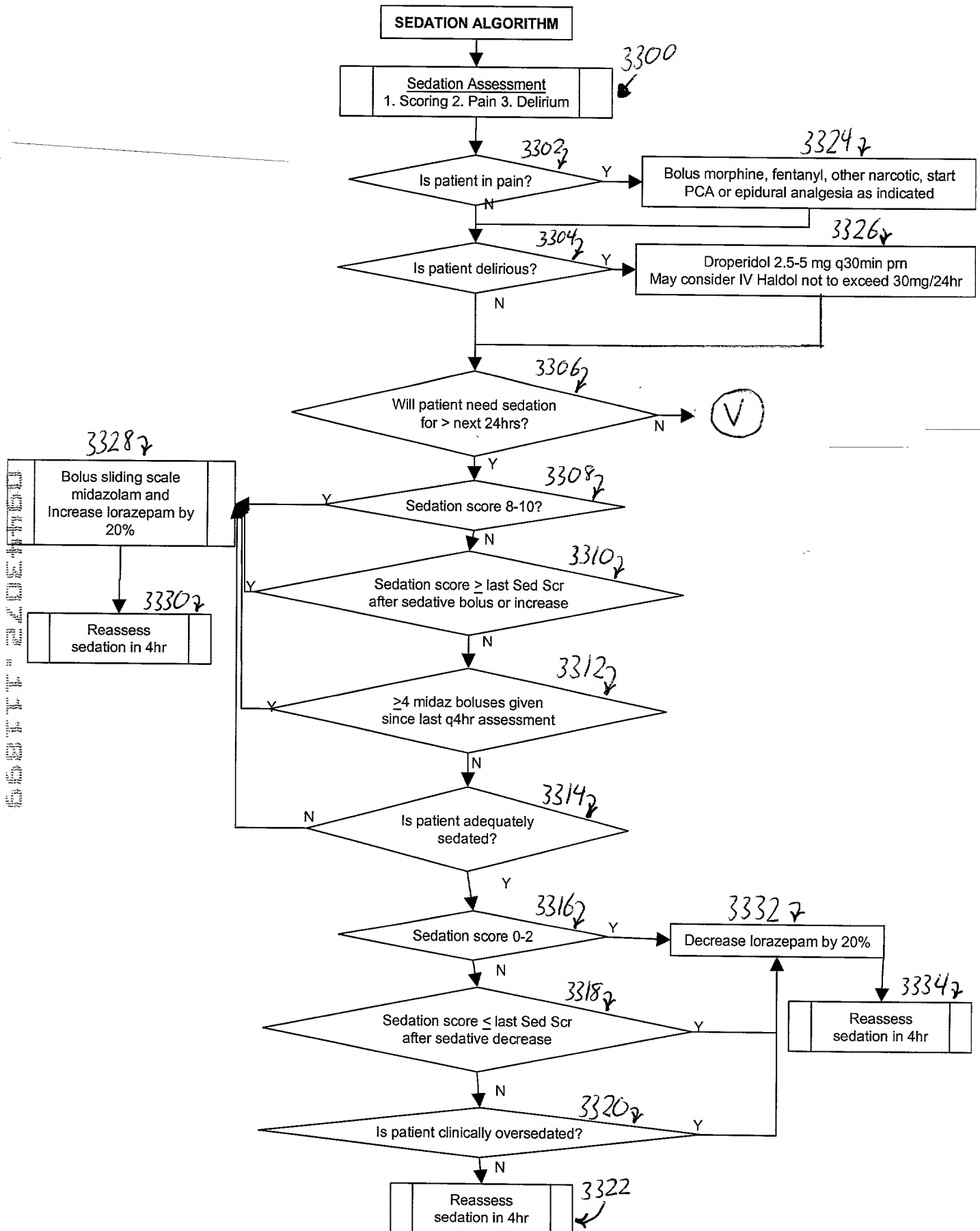
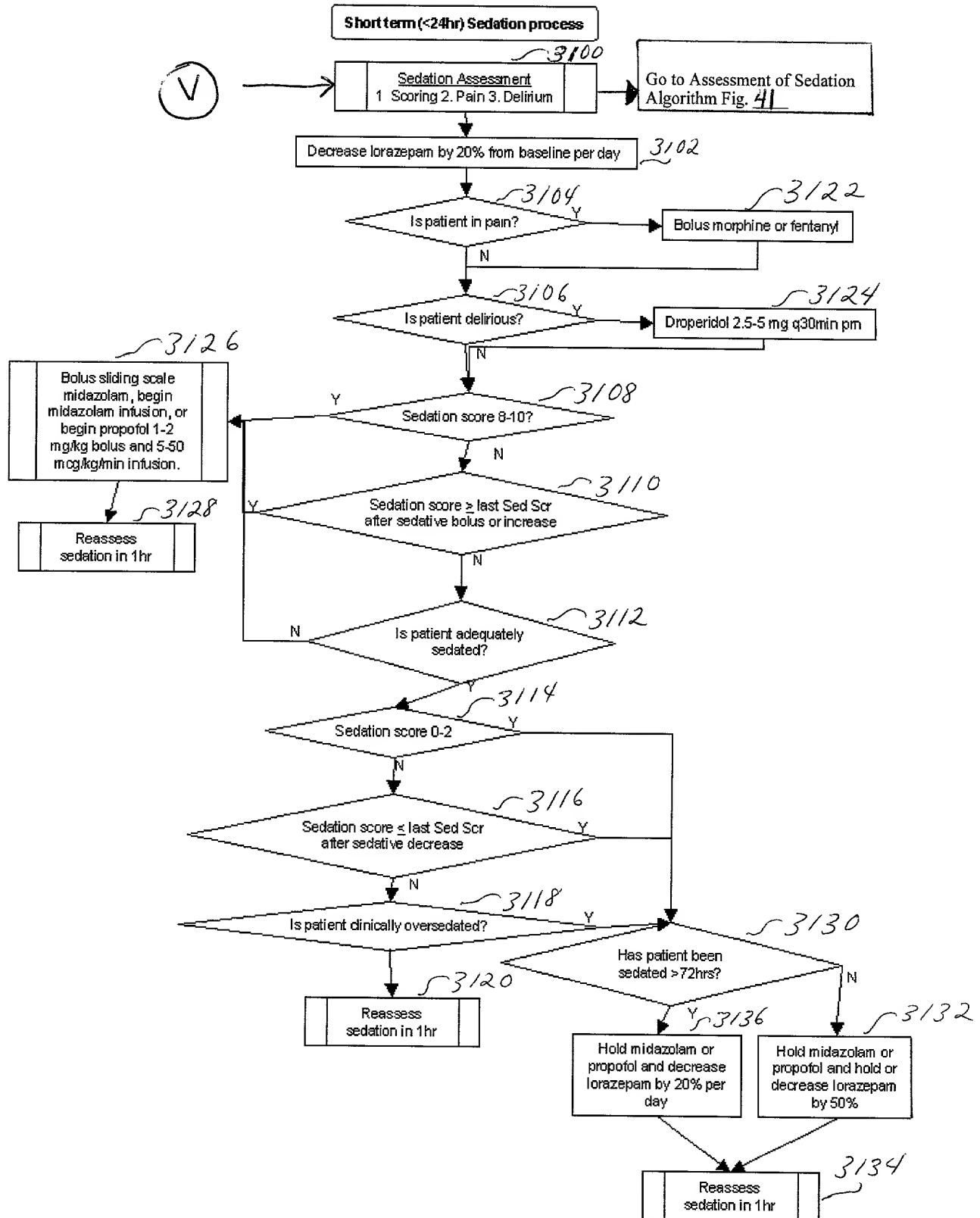


Figure 44





# Respiratory Isolation

Fig. 45

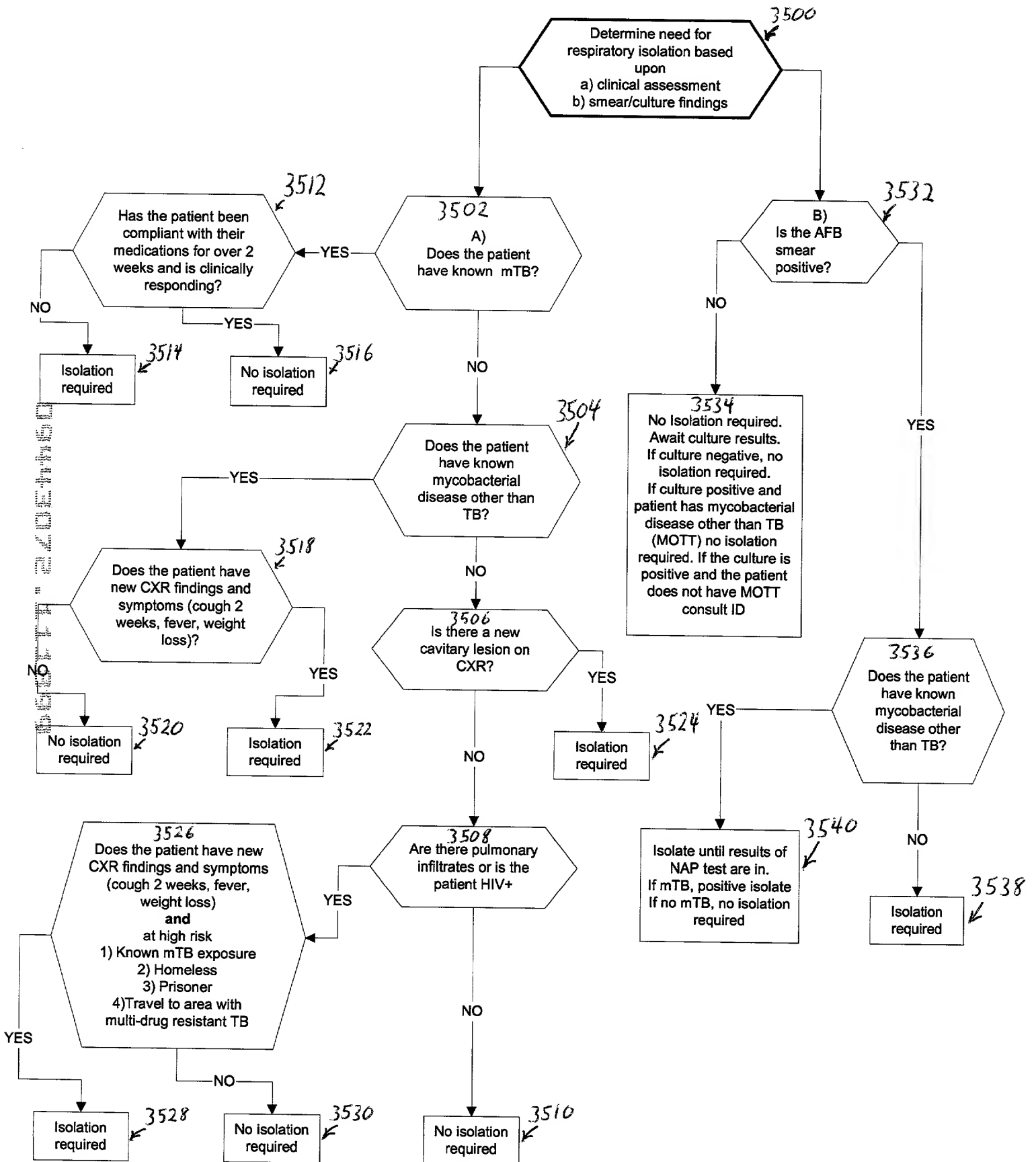
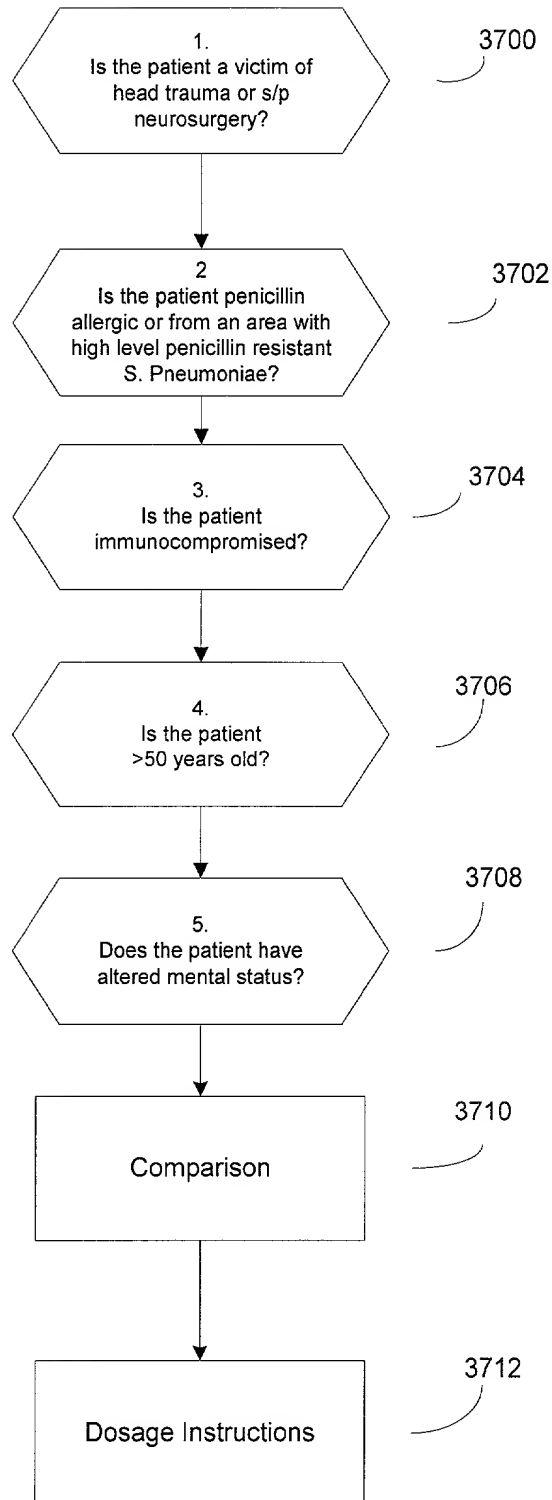


Figure 47

### Empiric Meningitis Treatment



# VENTILATOR WEANING

3800

<p><u>Daily Screen: perform between 0600 and 1000</u></p> <p>1. P/F &gt; 200 AND 2. PEEP ≤ 5 AND 3. Adequate cough with suctioning (or spontaneous) AND 4. No infusions of vasopressors (Dopamine ≤ 5 µg/kg/min) AND 5. No continuous infusions of sedatives or neuromuscular blocking agents</p>	
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3805

Rescreen next morning
-----------------------

3802

<p>All 5 screening criteria met?</p>
--------------------------------------

3804

<p><u>Rapid Shallow Breathing Test (Adapted from Tobin)</u></p> <p>1. Change mode to CPAP ≤ 5 (NO IMV and NO Pressure Support) 2. Give 1 minute to reach steady state 3. Measure <math>f/V_T</math> ratio where <math>f</math> = breaths per minute and <math>V_T</math> = tidal volume in Liters. If measured over a minute, <math>V_T</math> = minute ventilation/<math>f</math></p>
--

3806

<p>Rapid Shallow Breathing Test shows <math>f/V_T</math> ratio ≤ 105?</p>
---

3808

Rescreen next morning
-----------------------

3810

<p><u>Trial of Spontaneous Breathing</u></p> <p>Place on T-Piece or CPAP ≤ 5 (NO IMV or PS) for 2hr. Terminate if periodic assessment failed.</p>
---

3812

<p><u>Periodic Assessment</u></p> <p>Terminate weaning trial if:  <math>f &gt; 35</math> for 5 minutes OR  <math>SpO_2 &lt; 90\%</math> OR  <math>HR &gt; 140</math> OR  <math>HR &gt; 20\%</math> over or <math>&lt; 20\%</math> under baseline OR  <math>SBP &gt; 180</math> or <math>&lt; 90</math></p>
--

3814

<p>Successful 2hr trial of spontaneous breathing? AND  Able to defend airway (cough, no obstruction,  manageable secretions, requires suctioning less than  q2hr)? AND  No procedure planned next 24hr requiring patient to  be intubated?</p>
--

Progressive weaning
---------------------

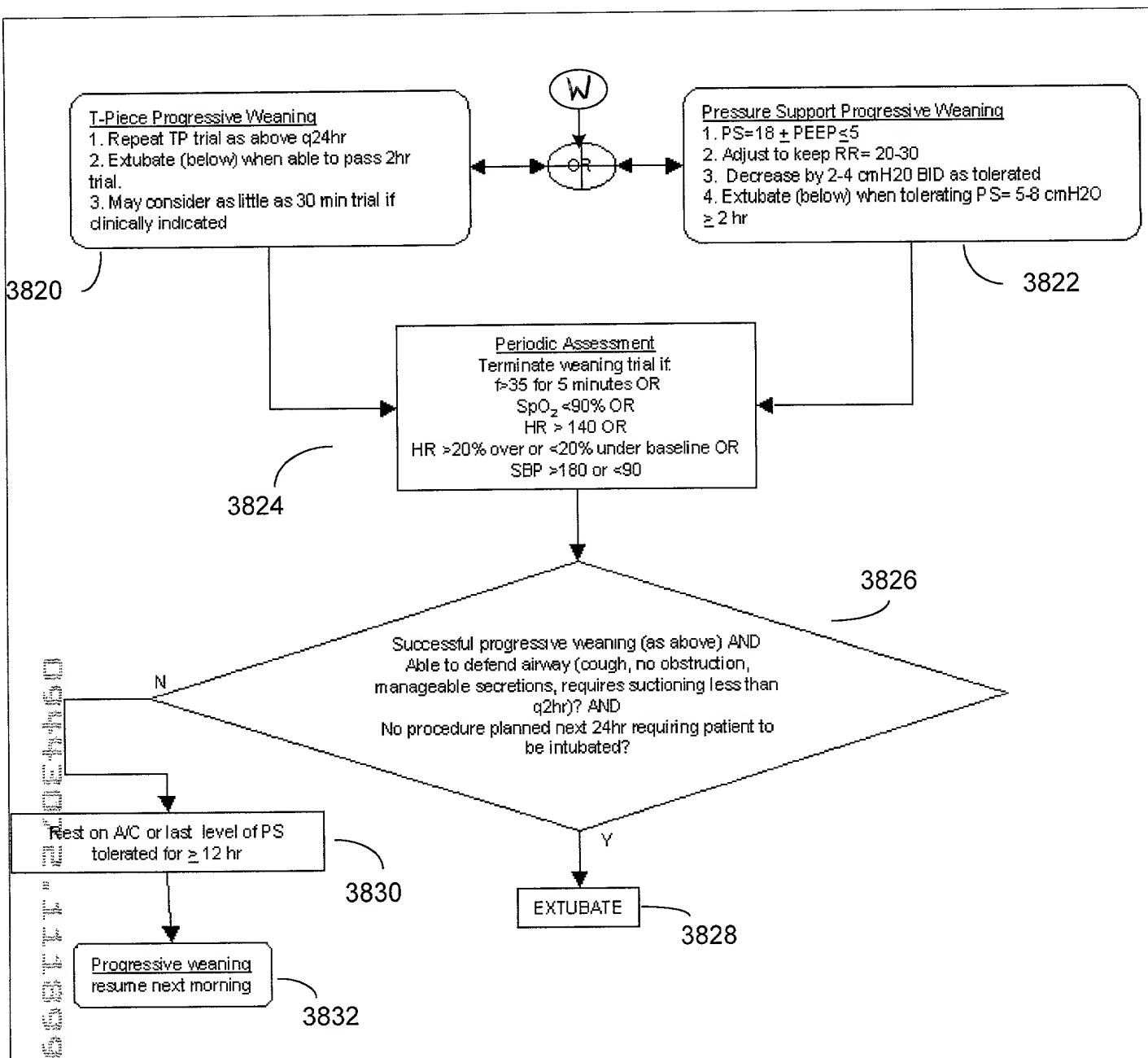
3816

EXTUBATE
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3818



Figure 48



**Figure 48A**

Figure 49

## Warfarin Dosing Algorithm

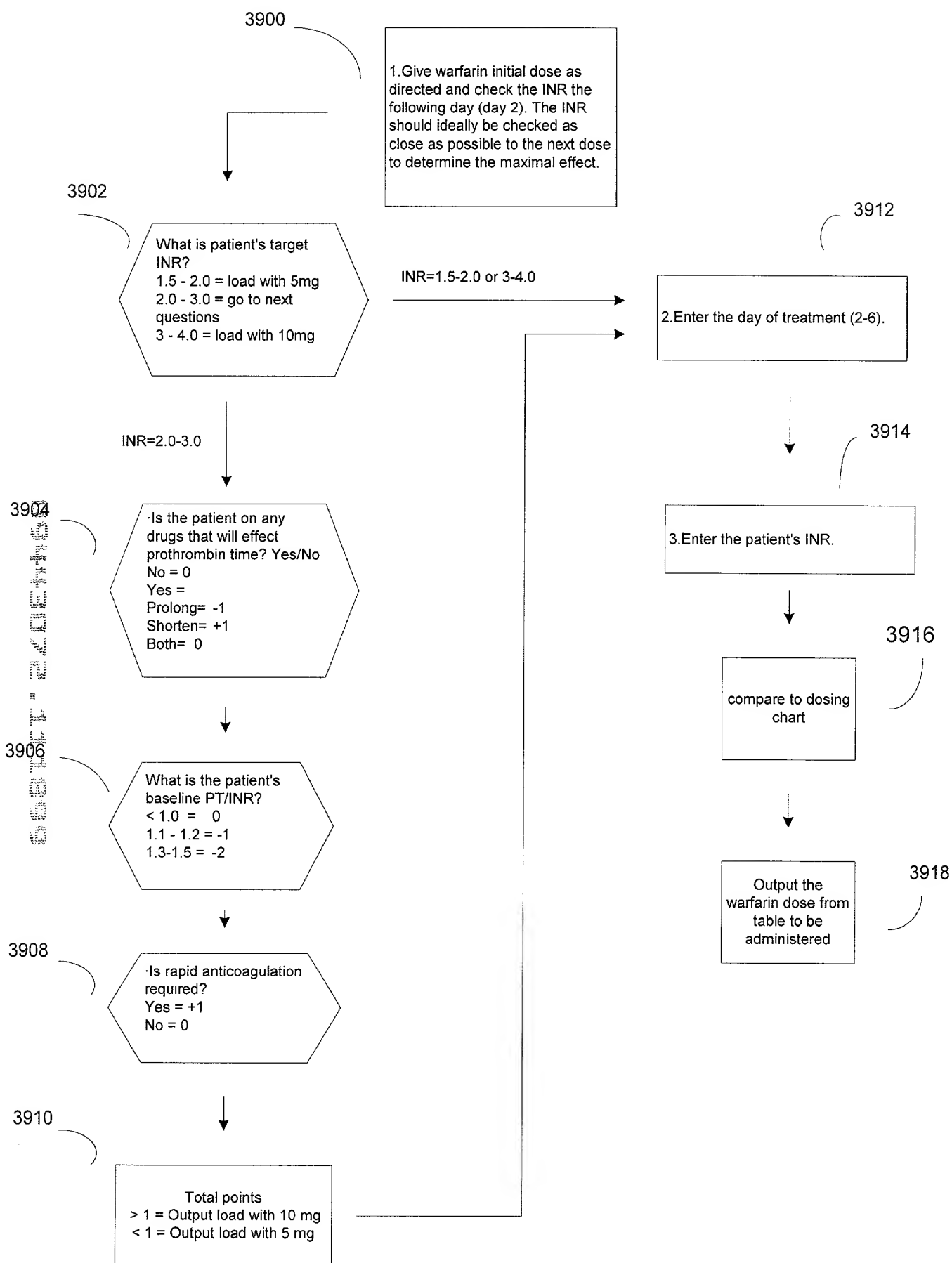
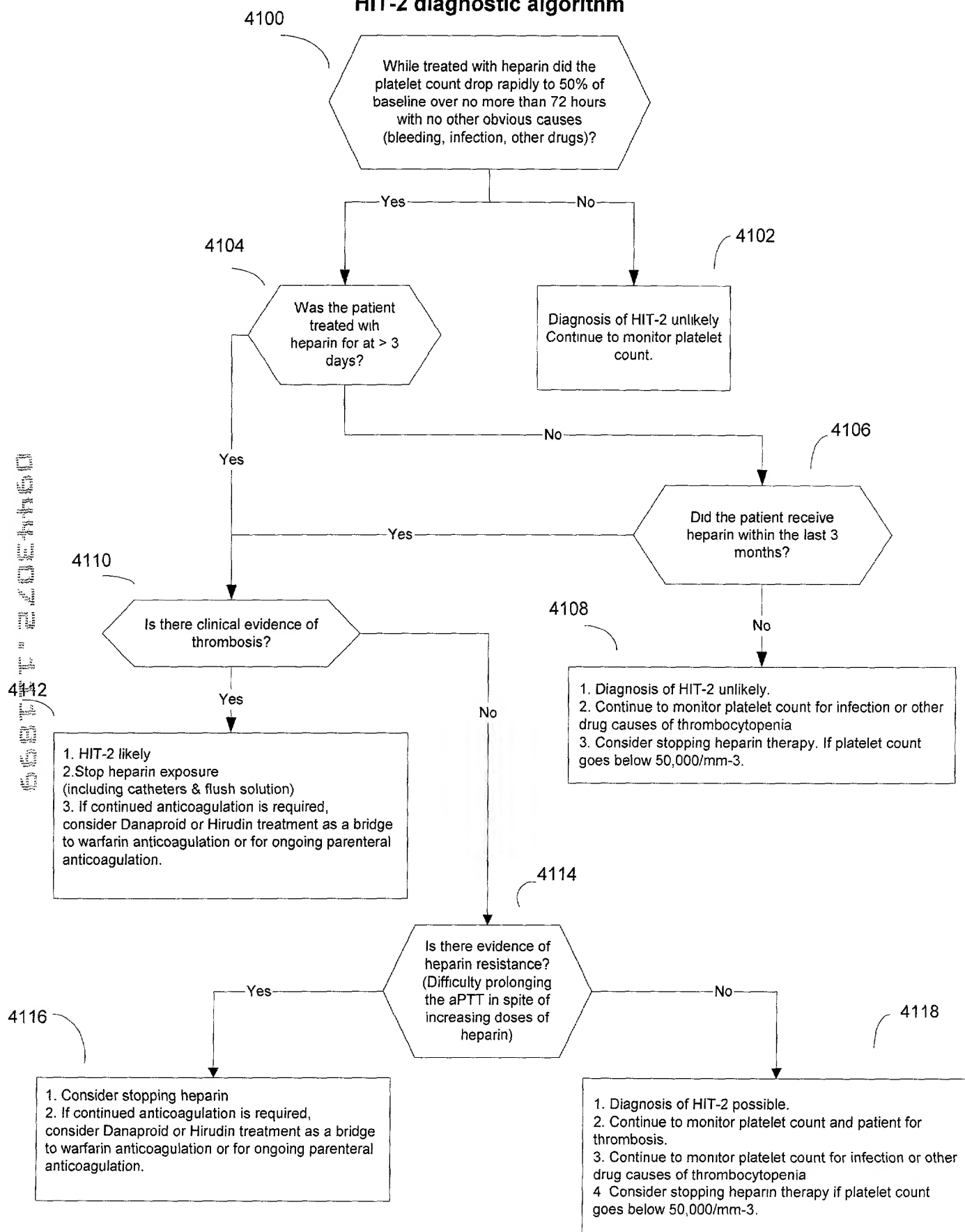


Figure 51

# HIT-2 diagnostic algorithm



**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Application of Brian A. Rosenfeld, M.D. and Michael Breslow

Serial No.: Not Yet Issued

Group Art Unit:

Filed: HERewith

Examiner:

FOR: **SYSTEM AND METHOD FOR PROVIDING CONTINUOUS, EXPERT NETWORK  
CRITICAL CARE SERVICES FROM A REMOTE LOCATION(S).**

**DECLARATION AND POWER OF ATTORNEY FOR PATENT APPLICATION**

As below inventors, we hereby declare that:

Our residence, post office address and citizenship are as stated below next to our names.

We believe we are the original, first and joint inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled **SYSTEM AND METHOD FOR PROVIDING CONTINUOUS, EXPERT NETWORK CRITICAL CARE SERVICES FROM A REMOTE LOCATION(S)**, the specification of which is attached hereto.

We hereby state that we have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

We acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, § 1.56(a).

We hereby appoint the following attorney(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

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0544333-44896

We declare that all statements made herein of our own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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